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**APOMORPHINE-CONTAINING DOSAGE FORM FOR  
AMELIORATING MALE ERECTILE DYSFUNCTION**

Cross-Reference to Related Applications

5           This application is a continuation-in-part of  
our co-pending application U.S. Serial No. 09/606,919,  
filed on June 29, 2000 and now U.S. Patent No.  
6,306,437, which is a continuation of U.S. Serial No.  
09/102,406, filed on June 22, 1998 and now U.S. Patent  
10       No. 6,121,276, which is a continuation-in-part of U.S.  
Serial No. 08/546,498 filed on October 20, 1995 and now  
U.S. Patent No. 5,770,606, which in turn is a  
continuation-in-part of U.S. Serial No. 08/231,250,  
filed on April 22, 1994, abandoned.

15       Field of the Invention

          This invention, in one aspect, relates to  
dosage forms and methods for ameliorating erectile  
dysfunction in psychogenic male patients. In another  
aspect this invention relates to diagnosis of erectile  
20       dysfunction. More particularly, this invention relates  
to the use of apomorphine-containing compositions for  
amelioration of erectile dysfunction in male patients  
and for diagnostic purposes.

Background of the Invention

25           A normal erection occurs as a result of a  
coordinated vascular event in the penis. This is  
usually triggered neurally and consists of vasodilation  
and smooth muscle relaxation in the penis and its  
supplying arterial vessels. Arterial inflow causes  
30       enlargement of the substance of the corpora cavernosa.  
Venous outflow is trapped by this enlargement,  
permitting sustained high blood pressures in the penis  
sufficient to cause rigidity. Muscles in the perineum  
also assist in creating and maintaining penile rigidity.  
35       Erection may be induced centrally in the nervous system

by sexual thoughts or fantasy, and is usually reinforced locally by reflex mechanisms. Erectile mechanics are substantially similar in the female for the clitoris.

Impotence or male erectile dysfunction is defined as the inability to achieve and sustain an erection sufficient for intercourse. Impotence in any given case can result from psychological disturbances (psychogenic), from physiological abnormalities in general (organic), from neurological disturbances (neurogenic), hormonal deficiencies (endocrine) or from a combination of the foregoing.

These descriptions are not exact, however. There is currently no standardized method of diagnosis or treatment. As used herein, psychogenic impotence is defined as functional impotence with no apparent overwhelming organic basis. It may be characterized by an ability to have an erection in response to some stimuli (e.g., masturbation, spontaneous nocturnal, spontaneous early morning, video erotica, etc.) but not others (e.g., partner or spousal attention).

Various methods for the treatment of impotence have been suggested, including external devices, for example, tourniquets (see U.S. Patent No. 2,818,855). In addition, penile implants, such as hinged or solid rods and inflatable, spring driven or hydraulic models, have been used for some time. The administration of erection effecting and enhancing drugs is taught in U.S. Patent No. 4,127,118 to LaTorre. That patent teaches a method of treating male impotence by injecting into the penis an appropriate vasodilator, in particular, an adrenergic blocking agent or a smooth muscle relaxant to effect and enhance an erection. More recently, U.S. Patent No. 4,801,587 to Voss et al. teaches the application of an ointment to relieve impotence. The

ointment consists of the vasodilators papaverine, hydralazine, sodium nitroprusside, phenoxybenzamine, or phentolamine and a carrier to assist absorption of the primary agent through the skin. U.S. Patent No.

5 5,256,652 to El-Rashidy teaches the use of an aqueous topical composition of a vasodilator such as papaverine together with hydroxypropyl- $\beta$ -cyclodextrin.

10 Recently the effect of apomorphine on penile tumescence in male patients afflicted with psychogenic impotence has been studied. These studies show that while apomorphine can indeed induce an erection in a psychogenic male patient, the apomorphine dose required to achieve a significant erectile response is usually accompanied by nausea or other serious undesirable side effects such as hypertension, flushing and diaphoresis. 15 The specific mechanisms by which apomorphine acts to produce an erectile response in a human patient are not yet completely understood, however.

20 Moreover, apomorphine has been shown to have very poor oral bioavailability. See, for example, Baldessarini et al., in Gessa et al., eds., Apomorphine and Other Dopaminomimetics, Basic Pharmacology, Vol. 1, Raven Press, N.Y. (1981), pp. 219-228.

25 Thus the search is continuing for an effective treatment of psychogenic impotence in male patients as well as for diagnostic methods that can identify such patients. It has now been found that certain delivery systems for apomorphine can provide a practical therapeutic and/or diagnostic "window" while reducing 30 the likelihood of undesirable side effects. It has also been found that nausea side effects associated with the use of apomorphine can be substantially reduced by the pre-administration or co-administration of an antiemetic agent.

Summary of the Invention

It has now been found that, for an optimal  
erectile response, steady state circulating serum and  
mid-brain tissue levels of apomorphine are to be  
5 maintained within a relatively closely defined range.

Nasal spray apomorphine dosage forms, usually  
containing about 1 to about 3 milligrams of apomorphine,  
are effective in male patients suffering from erectile  
dysfunction for the induction and maintenance of an  
10 erection sufficient for intercourse (i.e., vaginal  
penetration) without nausea or other undesirable side  
effects. The apomorphine is administered intranasally,  
preferably about 15 to about 20 minutes prior to sexual  
activity, and so as to maintain a predetermined  
15 circulating serum levels and mid-brain tissue levels of  
apomorphine during the period of sexual activity  
sufficient to induce an erection adequate for vaginal  
penetration but less than the amount that induces  
nausea. The plasma concentration of apomorphine should  
20 be maintained at no more than about 5.5 nanograms per  
milliliter, preferably about 0.3 to about 4 nanograms  
per milliliter, and more preferably about 1 to about 2  
nanograms per milliliter.

The foregoing intranasal apomorphine dosage  
25 forms are also suitable for screening patients  
complaining of erectile dysfunction.

The nausea side effect associated with the use  
of apomorphine can be substantially reduced by  
administration of an antiemetic agent together with the  
30 apomorphine. Specifically, a method suitable for  
treating erectile dysfunction in a male patient  
comprises administering to the patient prior to sexual  
activity, an antiemetic agent in an amount sufficient to  
substantially reduce nausea associated with use of

apomorphine, and apomorphine in an amount sufficient to induce and maintain an erection adequate for vaginal penetration.

The antiemetic agent is preferably co-administered with the apomorphine in a single nasal spray dosage unit. Separate dosage units with differing delivery routes are also suitable for practicing the present invention, however. For example, the antiemetic agent and apomorphine may be administered to the patient sequentially by first administering a composition comprising an antiemetic agent by any desired route of administration and thereafter a nasal spray composition comprising apomorphine.

A nasal spray dosage form for administering the antiemetic-apomorphine combination comprises apomorphine, an antioxidant, and optionally an antiemetic agent, in an aqueous pharmacologically acceptable liquid vehicle. The preferred anti-emetic agent is domperidone.

#### Brief Description of the Drawings

In the drawings,

FIGURE 1 is a graphical representation of mean erectile function, expressed as RIGISCAN™ monitor value, as a function of apomorphine dose;

FIGURE 2 is a bar graph depicting the percent successful erectile function for placebo, 3-milligram apomorphine dose, and 4-milligram apomorphine dose under erotic and neutral conditions;

FIGURE 3 is a bar graph presenting yet another comparison of erectile function noted in Pilot Study #4 in terms of RIGISCAN™ monitor score versus placebo, 3 milligrams of apomorphine and 4 milligrams of apomorphine under erotic and neutral conditions;

FIGURE 4 is a graphical representation of a comparison of the plasma concentration time profiles of apomorphine after intravenous administration at a dose of 1 mg (open circles, n=7), sublingual administration at a dose of 4 mg (open squares, n=4) and sublingual administration at a dose of 8 mg (half-filled squares, n=4);

FIGURE 5 is a graph of the dissolution pattern of apomorphine and the antiemetic agent nicotine for the tablets of Example 1;

FIGURE 6 is a graph of the dissolution pattern of apomorphine and the antiemetic agent nicotine for the tablets of Example 2;

FIGURE 7 is a graph of the dissolution pattern of apomorphine and the antiemetic agent nicotine for the layered tablets of Example 3;

FIGURE 8 is a graph of the dissolution pattern of apomorphine and the antiemetic agent prochlorperazine for the tablets of Example 4;

FIGURE 9 is a graph of the dissolution pattern of apomorphine and the antiemetic agent prochlorperazine for the layered tablets of Example 5;

FIGURE 10 is a graph of the dissolution pattern of apomorphine and the antiemetic agent prochlorperazine for the layered tablets of Example 6;

FIGURE 11 is a graph of the dissolution pattern of apomorphine and the antiemetic agent prochlorperazine for the layered tablets of Example 7;

FIGURE 12 is a graph of the dissolution of apomorphine for a sublingual apomorphine tablet as discussed in Example 8; and

FIGURE 13 is a graph comparing the dissolution pattern for the layered tablets of Example 7 with the

dissolution of apomorphine for a commercially available soluble apomorphine tablet.

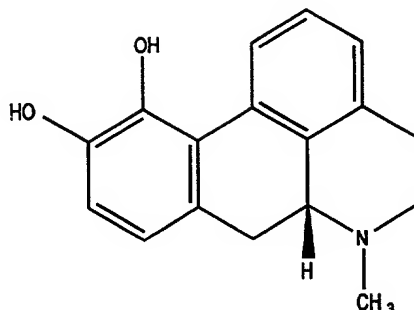
Detailed Description of Preferred Embodiments

5 Apomorphine is a dopamine receptor agonist  
that has a recognized use as an emetic when administered  
subcutaneously in about a 5-milligram dose. For the  
purposes of the present invention, apomorphine or a  
similarly acting dopamine receptor agonist is  
administered in an amount sufficient to excite cells in  
10 the mid-brain region of the patient but with minimal  
side effects. This cell excitation is believed to be  
part of a cascade of stimulation that is likely to  
include neurotransmission with serotonin and oxytocin.

The dopamine receptors in the mid-brain region  
15 of a patient can be stimulated to a degree sufficient to  
cause an erection by the sublingual administration of  
apomorphine so as to maintain a plasma concentration of  
apomorphine of no more than about 5.5 nanograms per  
milliliter (5.5 ng/ml). The sublingual administration  
20 usually takes place over a time period in the range of  
about 2 to about 10 minutes, or longer. The amount of  
apomorphine administered sublingually over this time  
period preferably is in the range of about 25 micrograms  
per kilogram ( $\mu\text{g/kg}$ ) of body weight to about 60  $\mu\text{g/kg}$  of  
25 body weight.

The apomorphine is administered preferably  
about 15 to about 20 minutes prior to anticipated sexual  
activity.

Apomorphine can be represented by the formula



and exists in a free base form or as an acid addition  
salt. For the purposes of the present invention  
5 apomorphine hydrochloride is preferred; however, other  
pharmacologically acceptable moieties thereof can be  
utilized as well. The term "apomorphine" as used herein  
includes the free base form of this compound as well as  
the pharmacologically acceptable acid addition salts  
10 thereof. In addition to the hydrochloride salt, other  
acceptable acid addition salts are the hydrobromide, the  
hydroiodide, the bisulfate, the phosphate, the acid  
phosphate, the lactate, the citrate, the tartarate, the  
salicylate, the succinate, the maleate, the gluconate,  
15 and the like.

Illustrative preferred sublingual dosage forms  
are set forth in Table I, below.



TABLE I

150-Milligram Apomorphine Hydrochloride Sublingual Tablets

	<u>3-mg Tablet</u>	
5	Apomorphine Hydrochloride	2.00 wt-%
	Mannitol	66.67 wt-%
	Ascorbic Acid	3.33 wt-%
	Citric Acid	2.00 wt-%
	Avicel PH102	15.00 wt-%
	Methocel E4M	10.00 wt-%
10	Aspartame	0.67 wt-%
	Magnesium Stearate	0.33 wt-%
	<u>4-mg Tablet</u>	
	Apomorphine Hydrochloride	2.66 wt-%
15	Mannitol	66.00 wt-%
	Ascorbic Acid	3.33 wt-%
	Citric Acid	2.00 wt-%
	Avicel PH102	15.00 wt-%
	Methocel E4M	10.00 wt-%
	Aspartame	0.67 wt-%
20	Magnesium Stearate	0.33 wt-%
	<u>5-mg Tablet</u>	
	Apomorphine Hydrochloride	3.33 wt-%
	Mannitol	65.34 wt-%
25	Ascorbic Acid	3.33 wt-%
	Citric Acid	2.00 wt-%
	Avicel PH102	15.00 wt-%
	Methocel E4M	10.00 wt-%
	Aspartame	0.67 wt-%
30	Magnesium Stearate	0.33 wt-%

If desired, and in order to facilitate absorption and thus bioavailability, the presently contemplated dosage forms can also contain, in addition to tabletting excipients,  $\beta$ -cyclodextrin or a  $\beta$ -cyclodextrin derivative such as hydroxypropyl- $\beta$ -cyclodextrin (HPBCD). Illustrative dosage forms containing HPBCD are shown in Tables II and III, below.

TABLE II

**Apomorphine Hydrochloride Sublingual Tablets  
With Hydroxypropyl- $\beta$ -Cyclodextrin**

	<u>mg/Tab</u>
5	
Apomorphine Hydrochloride	4.0
HPBCD	5.0
Ascorbic Acid	10.0
PEG8000	39.5
10 Mannitol	39.5
Aspartame	<u>2.0</u>
TOTAL	100.0

TABLE III

**Apomorphine Hydrochloride Sublingual  
Tablets With  $\beta$ -Cyclodextrin**

	<u>mg/Tab</u>
15	
Apomorphine Hydrochloride	5.0
20 $\beta$ -Cyclodextrin	20.0
Ascorbic Acid	5.0
Mannitol	68.9
Magnesium Stearate	1.0
D&C Yellow 10 Aluminum Lake	<u>0.1</u>
25 TOTAL	100.0

Apomorphine may be included in a nasal spray composition comprising apomorphine and a physiologically tolerable diluent. The present invention includes apomorphine and salts thereof formulated into sprayable compositions together with one or more non-toxic, physiologically tolerable or acceptable diluent, carrier, adjuvant or vehicle collectively referred to herein as a diluent for intranasal delivery.

Nasal spray dosage forms preferably comprise an aqueous apomorphine solution packaged in a nasal spray device, e.g., a pump driven spray device or the

like, capable of delivering a spray aliquot that contains apomorphine in an amount sufficient to induce an erection adequate for vaginal penetration but less than an amount that induces substantial nausea. The dosage form preferably is acidic, usually in the pH range of about 3 to 4. Antioxidants and preservatives can be included in the spray dosage form if desired.

These nasal spray compositions can also contain adjuvants such as preservatives and/or wetting, emulsifying, and dispensing agents. Inhibition of microorganisms can be achieved by various antibacterial and antifungal agents, for example, the parabens, chlorobutanol, phenol, sorbic acid, and the like. It may also be desirable to include isotonic agents, for example sugars and sodium chloride, among others.

Preferred compositions for intranasal delivery contain a stabilizer and a surfactant. Among the pharmaceutically acceptable surfactants are polyoxyethylene castor oil derivatives, such as polyoxyethylene-glycerol-triricinoleate, also known as polyoxoyl 35 castor oil (CREMOPHOR EL), or poloxyl 40 hydrogenated castor oil (CREMOPHOR RH40) both available from BASF Corp.; mono-fatty acid esters of polyoxyethylene (20) sorbitan, such as polyoxyethylene (20) sorbitan monolaurate (TWEEN 80), polyoxyethylene monostearate (TWEEN 60), polyoxyethylene (20) sorbitan monopalmitate (TWEEN 40), or polyoxyethylene 20 sorbitan monolaurate (TWEEN 20) all available from ICI Surfactants of Wilmington, DE); polyglyceryl esters, such as polyglyceryl oleate; and polyoxyethylated kernel oil (LABRAFIL, available from Gattefosse Corp.) Preferably, the surfactant will be between about 0.01% and 10% by weight of the pharmaceutical composition.

Among the pharmaceutically acceptable stabilizers are antioxidants such as sodium sulfite, sodium metabisulfite, sodium thiosulfate, sodium formaldehyde sulfoxylate, sulfur dioxide, ascorbic acid, isoascorbic acid, thioglycerol, thioglycolic acid, cysteine hydrochloride, acetyl cysteine, ascorbyl palmitate, hydroquinone, propyl gallate, nordihydroguaiaretic acid, butylated hydroxytoluene, butylated hydroxyanisole, alpha-tocopherol and lecithin. Preferably, the stabilizer is present in an amount in the range of about 0.01% and 5% by weight of the composition.

Chelating agents such as ethylene diamine tetraacetic acid, its derivatives and salts thereof, e.g, edetate disodium, as well as dihydroxyethyl glycine, citric acid, tartaric acid, and the like may also be utilized.

Illustrative nasal spray compositions are presented in Table IV, below.

**TABLE IV**  
**Nasal Spray Compositions**

	<u>A</u>	<u>B</u>	<u>C</u>	<u>D</u>	<u>E</u>
Apomorphine•HCl	0.6 g	0.5 g	0.05 g	0.005 g	1 g
Ascorbic acid	1 g	—	—	1 g	—
Edetate disodium	—	0.01 g	0.01 g	—	—
Sodium bisulfite	0.05 g	0.1 g	0.1 g	0.05 g	0.1 g
Sodium chloride	0.75 g	0.75 g	—	0.75 g	—
Hydrochloric acid (0.1N)	0.9 g	0.9 g	0.3 g	q.s.	q.s.
Distilled water, q.s.	1000 ml	1000 ml	1000 ml	1000 ml	1000 ml
pH	about 3-4	about 3.3	about 3.8	about 3-4	about 3-4

Optionally, the nasal spray compositions shown in Table IV can include an effective amount of an anti-emetic agent such as domperidone, nicotine, lobeline sulfate, and the like as discussed in greater detail hereinbelow.

5           The onset of nausea can be obviated or delayed by delivering apomorphine at a controlled rate so as to provide circulating serum levels and mid-brain tissue levels of apomorphine sufficient for an erection without inducing substantial nausea. When apomorphine is  
10 administered at or near the relatively higher amounts of the aforementioned dosage range, the likelihood of nausea onset can be reduced by concurrent administration of a ganglionic agent (inhibitor of ganglionic response) such as nicotine or lobeline sulfate. For this purpose,  
15 the weight ratio of apomorphine to ganglionic agent is in the range of about 300:1 to about 5:1.

          The preferred weight ratio necessarily varies according to the potency of the agent employed, however. When nicotine is used, the preferred weight ratio of  
20 apomorphine to nicotine is in the range of about 10 to about 1. With regard to specific drug loadings, sublingual dosage units for co-administration of nicotine and apomorphine preferably contain apomorphine in the range of about 1 to about 8 milligrams (mg) and  
25 nicotine in the range of about 0.25 to about 3 mg. A particularly preferred sublingual combination dosage unit contains apomorphine in the range of about 4 mg to about 8 mg, and nicotine in the range from about 0.75 mg to about 1.25 mg.

30           Nicotine and lobeline sulfate have been classified as ganglionic stimulating alkaloids. See, for example, Goodman, Louis S. and Alfred Gilman, eds., *The Pharmacological Basis of Therapeutics*, 5th Ed., MacMillan Publishing Co., New York, N.Y. (1975), pp.

567-569. For the purposes of the present invention, ganglionic stimulating alkaloids such as nicotine and lobeline sulfate serve as antiemetic agents.

Antiemetic agents are drugs that prevent or substantially reduce nausea and vomiting. As used herein, the terms "antiemetic agent" and "antinausea agent" are interchangeable and mean a pharmaceutically acceptable compound that substantially reduces nausea symptoms. As described below, antiemetics may be classified according to their structure or their mechanism of operation.

In addition to the ganglionic stimulating alkaloids discussed above, other antiemetic agents that can be used in conjunction with apomorphine are antidopaminergic agents such as metoclopramide, and the phenothiazines, e.g., chlorpromazine, prochlorperazine, pipamazine, thiethylperazine, oxypendyl hydrochloride, promazine, triflupromazine, propiomazine, acepromazine, acetophenazine, butaperazine, carphenazine, fluphenazine, perphenazine, thiopropazate, trifluoperazine, mesoridazine, piperacetazine, thioridazine, pipotiazine, pipotiazine palmitate, chlorprothixine, thiothixine, doxepin, loxapin, triflupromazine, methdilazine, trimeprazine, methotrimeprazine, and the like. Metoclopramide is a benzamide. Benzamides are a recognized group of antiemetics that are suitable for the present invention and include in addition to metoclopramide, trimethobenzamide and benzquinamid, as well as others. Also suitable are the serotonin (5-hydroxytryptamine or 5-HT) antagonists such as domperidone, ondansetron (commercially available as the hydrochloride salt under the designation Zofran®), and the like, the histamine antagonists such as buclizine hydrochloride, cyclizine

hydrochloride, dimenhydrinate (Dramamine), and the like, the parasympathetic depressants such as scopolamine, and the like, as well as other antiemetics such as metopimazine, trimethobenzamide, benzquinamine hydrochloride, diphenidol hydrochloride, and the like. Another suitable group of antiemetics are the meclizines which include, for example, meclizine, chlorcyclizine, cyclizine, and buclizine.

Accordingly, a composition aspect of the present invention provides a combination of apomorphine and an antiemetic agent which is a member of the group consisting of the phenothiazines, the benzamides, the meclizines, the serotonin antagonists, hydroxyzine, lobeline sulfate, dimenhydrinate, scopolamine, metopimazine, diphenidol hydrochloride, nicotine, and their acid addition salts.

Any pharmaceutically acceptable form of the antiemetic agents can be employed, i.e., the free base or a pharmaceutically acceptable salt thereof (e.g. cyclizine hydrochloride, cyclizine acetate, diphenhydramine hydrochloride, meclizine hydrochloride, etc.)

The nausea side effect associated with the use of apomorphine can be substantially reduced by administration of an antiemetic agent. Specifically, a method suitable for treating erectile dysfunction in a male patient comprises administering to the patient (prior to sexual activity) an antiemetic agent in an amount sufficient to substantially reduce nausea associated with use of apomorphine, and apomorphine in an amount sufficient to induce and maintain an erection adequate for vaginal penetration.

For treatments according to the present invention, an antiemetic agent may be co-administered

with apomorphine or may be administered concurrently or sequentially with apomorphine to substantially reduce the symptoms of nausea associated with the use of apomorphine. By the term "co-administration" is meant the administration of both apomorphine and an antiemetic agent to the patient in a single unit dosage form as, for example, in an aqueous spray. "Concurrent" administration denotes the substantially simultaneous administration of the two drugs in separate unit dosage forms, while "sequential" administration is the administration of separate dosage forms of the two drugs with one being administered at some time interval after the other. The co-administration of an antiemetic agent and apomorphine is preferred and allows for a higher dose of apomorphine with generally improved response and function.

When the antiemetic agent prochlorperazine hydrochloride is used, the preferred weight ratio of apomorphine hydrochloride to prochlorperazine hydrochloride is in the range of about 5 to about 0.25. The amount of prochlorperazine hydrochloride administered sublingually preferably is in the range of about 5 µg/kg of body weight to about 200 µg/kg of body weight.

Apomorphine with antiemetic-containing dosage forms including nicotine-containing dosage forms and domperidone-containing dosage forms are illustrated in Table V, below.



TABLE V

**Apomorphine Hydrochloride Sublingual Tablets  
Containing an Antiemetic Agent**

		<u>mg/Tab</u>
5	Apomorphine Hydrochloride	5.0
	Ascorbic Acid	5.0
	Mannitol	67.9
	Magnesium Stearate	1.0
10	Nicotine	1.0
	$\beta$ -Cyclodextrin	20.0
	D&C Yellow 10 Aluminum Lake	<u>0.1</u>
	TOTAL	100.0
15		<u>mg/Tab</u>
	Apomorphine Hydrochloride	5.0
	Ascorbic Acid	5.0
	Mannitol	58.9
20	Magnesium Stearate	1.0
	Domperidone	10.0
	$\beta$ -Cyclodextrin	20.0
	D&C Yellow 10 Aluminum Lake	<u>0.1</u>
	TOTAL	100.0
25		<u>mg/Tab</u>
	Apomorphine Hydrochloride	4.0
	Nicotine Base	1.0
30	Acesulfame-K	4.0
	Microcrystalline Cellulose	37.5
	Peppermint Flavor	2.5
	Chocolate Flavor	2.0
	Citric Acid	3.0
35	Hydroxypropylmethylcellulose	13.0
	Mannitol	80.0
	Magnesium Stearate	<u>3.0</u>
	TOTAL	150.0
40		

TABLE V-continued

		<u>mg/Tab</u>
5	Tablet core:	
	Apomorphine Hydrochloride	4.0
	Acesulfame-K	1.6
	Microcrystalline Cellulose	21.6
	Peppermint Flavor	1.0
10	Chocolate Flavor	0.8
	Citric Acid	1.2
	Hydroxypropylmethylcellulose	4.0
	Mannitol	24.6
	Magnesium Stearate	1.2
15	Tablet outer layer:	
	Nicotine Base	1.0
	Acesulfame-K	0.4
	Microcrystalline Cellulose	36.6
20	Mannitol	47.0
	Hydroxypropylmethylcellulose	4.0
	Magnesium Stearate	<u>1.0</u>
	TOTAL	150
		<u>mg/Tab</u>
25	Apomorphine Hydrochloride	4.0
	Prochlorperazine Hydrochloride	5.0
	Acesulfame-K	4.0
	Microcrystalline Cellulose	37.5
30	Peppermint Flavor	2.5
	Chocolate Flavor	2.0
	Citric Acid	3.0
	Hydroxypropylmethylcellulose	10.0
	Mannitol	68.0
35	Sodium Alginate	10.0
	Magnesium Stearate	<u>3.0</u>
	TOTAL	150.0

TABLE V-continued

		<u>mg/Tab</u>
5	Tablet core:	
	Apomorphine Hydrochloride	4.0
	Acesulfame-K	1.6
	Microcrystalline Cellulose	20.0
	Peppermint Flavor	1.0
10	Chocolate Flavor	0.8
	Citric Acid	1.2
	Hydroxypropylmethylcellulose	5.0
	Mannitol	20.2
	Sodium Alginate	5.0
15	Magnesium Stearate	1.2
	Tablet outer layer:	
	Prochlorperazine	5.0
	Acesulfame-K	0.4
20	Microcrystalline Cellulose	35.6
	Mannitol	46.0
	Hydroxypropylmethylcellulose	2.0
	Magnesium Stearate	<u>1.0</u>
	TOTAL	100.0
25		

For improved bioavailability, controlled release, and reliable dosage control, the apomorphine containing compositions of the present invention are preferably administered intranasally. The preferred  
5 nasal spray dosage forms contain about 1 to about 3.75 milligrams of apomorphine preferably about 1.25 to about 2.5 milligrams of apomorphine.

Antiemetic agents may also be delivered to patients using other conventional drug delivery methods,  
10 such as orally, intravenous injection, subcutaneous injection, suppository, or patch (e.g. buccal patch). In addition, patients may receive the apomorphine and the antiemetic agent via different delivery mechanisms. For example, the apomorphine may be delivered via a  
15 nasal spray, while the antiemetic agent is delivered orally.

When an antiemetic agent is used, the antiemetic agent is preferably made available before the apomorphine. This can be accomplished not only by  
20 administering the antiemetic agent before the apomorphine but also by employing a staggered release dosage form as described below. The present invention is also not limited to a particular sequence of administration or dosage form for the antiemetic agent  
25 and apomorphine. If desired, the antiemetic agent may be administered substantially concurrently (i.e., at the same time as) or even after the apomorphine. For example, a separate dosage form of an antiemetic agent can be made available to patients for use after  
30 administration of the apomorphine if nausea symptoms are encountered.

The antiemetic agent preferably is delivered to the patient with the apomorphine via a single dosage unit. Provided for this purpose, a sublingual tablet

comprises apomorphine, an antiemetic agent, an osmotic agent, and a swellable hydrophilic carrier. A preferred swellable hydrophilic carrier is microcrystalline cellulose. Other suitable swellable hydrophilic carriers for the present purposes are ethyl cellulose, microcrystalline cellulose, cross-linked polyvinyl pyrrolidone, dicalcium phosphate, calcium carbonate and silica.

Suitable osmotic agents include monosaccharide and disaccharide sugars, such as glucose, fructose, mannitol, sorbitol, lactose, and sucrose. Glycerin or urea may also be used. Organic and inorganic salts, such as sodium chloride, potassium chloride and water soluble polyelectrolytes, are also suitable as osmotic agents. A preferred osmotic agent is mannitol. Preferred embodiments of a sublingual tablet according to the present invention also contain a lubricant such as magnesium stearate.

One aspect of the present invention provides an apomorphine/antiemetic combination formulated into a dosage unit that provides a staggered release of antiemetic agent and apomorphine. Specifically, a dosage unit includes an antiemetic agent as a relatively faster release component and apomorphine as a component released after release of the antiemetic agent has begun. Defined in terms of release rate, one preferred apomorphine/antiemetic dosage unit obtains 50 percent release of the antiemetic agent at least 5 minutes before obtaining 50 percent release of the apomorphine.

For this staggered release purpose, the present invention further provides a layered tablet that comprises a core layer containing apomorphine and an outer layer containing an antiemetic agent. Table V

(above) as well as the Examples below contain illustrative compositions for layered tablets.

5 Illustrative preferred sublingual dosage forms for apomorphine/antiemetic combinations are set forth in the Examples 1-7.

10 The present invention is illustrated further by the following studies which were focused on two specific objectives. The first was to determine whether, relative to placebo response, patients who presented with "psychogenic" impotence (i.e., patients who were still capable of achieving erections) demonstrated improved erectile function and/or enhanced sexual desire post-dosing with sublingual apomorphine (APO). The second objective was to determine what  
15 dose(s) of various forms of sublingual APO are effective in this group of patients for inducing an erection that is sufficient for vaginal penetration.

20 Participating patients were selected from among those that initially presented with the complaint of impotence. These patients underwent a thorough urological assessment by a urologist as well as an assessment by a psychiatrist. Diagnostic testing for erectile difficulties was extensive and included the following: biochemical profile, nocturnal penile  
25 tumescence (NPT) monitoring, doppler flow studies, biothesiometry, corporal calibration testing with an intracorporal injection of triple therapy and dynamic cavernosometry. These tests were used to rule out any arterial, venous or peripheral neural causality of  
30 impotence. Any patients with abnormalities in any of these three areas were excluded from entry to the trials. The inclusion/exclusion criteria for all four pilot studies are set forth in Table VI, below. Patients who met all criteria were diagnosed as having

impotence primarily of a psychogenic origin. If there were no known medical contraindications to the use of a dopaminergic medication they were offered entry into an APO trial.

5                   Instructions were given regarding the protocol by the research clinician, and an informed consent was obtained. Patients were advised that they were free to withdraw from the trial at any time without penalty or prejudice. They were tested on at least three separate  
10                   days at three separate doses (placebo and two active medication doses) with an interval of no less than three days between. The experimental scheme described below was used in all four pilot studies.

                  Patients were seated in a comfortable chair  
15                   and a RIGISCAN<sup>TM</sup> ambulatory tumescence monitor (Dacomed Corp., Minneapolis, Minnesota) was placed on the patient and the computer was set in the real time monitoring mode. Blood pressure and heart rate were recorded pre-dosing with APO or placebo and at the end of the testing  
20                   session. Visual analogue scales (VAS) were completed by the patient pre-dosing as well as post-dosing (at the end of the testing session). These scales reflected the patient's sense of well being, level of sedation, tranquilization, anxiousness, arousal and any changes in  
25                   yawning behavior. In a single-blind fashion, apomorphine or placebo was administered to the patient sublingually. Doses of active medication varied on the formulation of the apomorphine administered (liquid or tablet). Because of the possibility of nausea and the  
30                   tolerance to this effect that prior dosing conveys, the patient was given increasing doses at each testing. However, the patient was unaware of the dose that he was receiving (single-blind). Patients were instructed not

to swallow the medication, but to keep it under their tongue and allow it to be absorbed there.

Symptoms as they were volunteered were recorded by the research clinician. If the patient complained of nausea or felt unwell in any way he was asked if he wanted to abort the trial. If the trial was aborted, the patient was given Gravol 50 mg p.o. at that time. The patient was monitored by the research clinician until these side-effects had subsided. He was asked to return the following week for retesting at the same dose and was instructed to begin treatment with Domperidone 10 m.g. p.o. TID the day before and morning of his next session.

Patients not experiencing nausea or any other significant adverse effects within fifteen minutes post-dosing with APO or placebo viewed segments of standardized erotic videos to provide sexual stimulation. The following sequence of videos was viewed: a ten minute erotic video, a neutral video lasting between five and ten minutes in duration and finally another ten minute erotic video. The duration of the testing session for each dose level lasted between 45 and 60 minutes. After determining the most effective dose of apomorphine for the patient, he was then offered APO for domestic trial at that dose.

#### Results of Pilot Studies 1 to 4

The frequency and the magnitude of erectile responses were documented with each dose of apomorphine or placebo. Data obtained from the RIGISCAN™ monitor was downloaded and each session was scanned. Erection responses were then scored for rigidity (%) and tumescence (cm.) at both the tip and base of the penis and an overall score was given that corresponded to these parameters during the viewing of both erotic and



neutral video segments (see Table VII, below). A score of less than 16 indicated erectile dysfunction and a poor response to apomorphine at that dose.

5 Visual analogue scales (See Table X) were compared both pre- and post-dosing, and examined for changes in feeling of well being, levels of arousal, anxiousness, sedation/tranquilization and yawning behavior. Blood pressure and heart rate were also compared pre- and post-dosing.

10 Effects of apomorphine that were both reported to and observed by the research clinician were grouped into two categories: Adverse Effects (i.e., flushing, diaphoresis, nausea, vomiting, changes in blood pressure or heart rate) or Primary Effects (i.e., yawning and erections).

15 Each pilot study was reviewed under the categories mentioned above.

#### Pilot Study #1

20 The initial formulation evaluated was liquid apomorphine administered via sublingual route. APO was prepared by a clinic pharmacist and dissolved in a solution of sodium metabisulfite and ethylenediamine tetraacetic acid (EDTA). The final concentration was 100 mg/ml. Patients were tested on three separate occasions at three separate doses (placebo; 10 mg; 20 mg)

25 Twelve patients entered into this trial. All patients had reported erectile dysfunction greater than 1 year in duration. The age range in this group was 30 from 38 to 60 years. One patient withdrew after placebo and another withdrew after adverse effects at the 20 mg dose. That left a total evaluable group of ten. All ten patients had previously received yohimbine HCl for erectile dysfunction. Eight had failed a trial of

yohimbine HCl. Of this group of eight, 6 were successful with apomorphine.

Seven (70%) were successes (score of no less than 16 on both neutral and erotic video segments; Table VII) and three (30%) were categorized as failures with apomorphine. Six out of the seven successful patients continued on with a domestic trial of apomorphine at the dose that gave them the best response during testing. Three required treatment with Domperidone the day before and morning of apomorphine usage. The range of domestic use varied from two to seven months.

Analysis of visual analogue scales pre- and post-dosing with apomorphine indicated the following. At the end of the session patients were relaxed but not sedated. There was no evidence of arousal or anxiousness. Yawning behavior changes were evident on these scales with the incidence of yawning increasing between 15 and fifty minutes post-dosing and with each increase in dosing. Each patient experienced between two to five yawns per session. These changes were not evident with placebo.

The primary effect of yawning was both reported by patients and observed at both 10 mg and 20 mg doses. No yawning was reported with placebo. Adverse effects were reported at both dose levels. Two patients who did not experience nausea or diaphoresis were researched for similarities in their patient profiles but none were found. Anywhere from ten to fifteen minutes post-dosing the other eight patients developed sudden onset of various levels of nausea (and in one instance vomiting), diaphoresis, dizziness, double or blurred vision, decrease in both blood pressure and heart rate and pale or ashen coloring. Side effects varied from being transient and brief to

lasting as long as from 30 to 40 minutes. One patient reported a stuffy nose starting approximately 30 minutes post-dosing and lasting for approximately 10 minutes. No adverse effects were reported post placebo dosing.

5           The foregoing Pilot Study leads to the following conclusions:

1. Apomorphine is effective in inducing erectile episodes without increasing libido in the "psychogenically" impotent male.

10           2. Both 10 mg and 20 mg doses produce erectile responses.

3. Both doses produced adverse effects (nausea, vomiting, diaphoresis, etc.) that would be unacceptable to patients and their partners, however. These effects can be counteracted with the use of Domperidone.

15           Pilot Study #2

20           The first sublingual tablet formulations evaluated were 2.5 and 5 mg. Patients were tested on three separate occasions at three separate doses (placebo; 2.5 mg, 5 mg).

25           A total of eight patients entered into this trial. All patients reported erectile difficulties for more than two years. The age range was from 38 to 62 years. All had failed a trial of yohimbine HCl. One patient withdrew from the trial after experiencing adverse effects at the 5 mg dose. That left a total of seven evaluable patients.

30           Two (29%) were successes (score of no less than 16; Table VII) and five (71%) were failures during lab testing. The two successful patients went onto a domestic trial of apomorphine at the 2.5 mg dose which was the most effective and did not produce adverse

effects. Both patients used apomorphine at home for no less than two months with satisfactory results.

Analysis of visual analogue scales pre- and post-dosing with apomorphine indicated the same trends as with the liquid apomorphine preparation. Patients were relaxed but not sedated. No evidence of arousal or anxiousness was noted.

The primary effect of yawning was both reported by patients and observed at both 2.5 mg and 5 mg doses. The incidence of yawning increased between fifteen and forty minutes post-dosing. At the 2.5 mg dose all patients who failed testing had only one or two yawns per session. The 5 mg dose not only produced adverse effects (nausea, diaphoresis, dizziness, blurred vision, facial flushing, drop in both heart rate and blood pressure) but also increased yawning responses to three to five times per session. The two successful patients experienced three to five yawns at both the 2.5 mg and 5 mg doses. These changes were not evident with placebo.

At the end of Pilot Study #2 the following conclusions were made:

1. There appears to be a correlation between the effectiveness of the dose and yawning response (poor responders experience less yawning).

2. Both 2.5 and 5 mg doses produced erectile responses in some patients. The apparent 28% success rate was because of lab use only (failures were not given drug to take home) and lack of available intermediate doses.

3. In some instances the 5 mg dose can produce adverse effects (i.e., nausea, diaphoresis, etc.) that may be unacceptable to patients and their partners. These effects can be counteracted with the

administration of Domperidone or nicotine (e.g., by smoking).

4. The sublingual tablets were easy to administer and dissolved within five minutes.

5 Pilot Study #3

Apomorphine was evaluated as an aqueous intranasal spray (1.25 mg per puff). The first patient was an anxious, 53 year old male who had been experiencing erectile dysfunction for two years. This patient had previously failed a trial of yohimbine.

He was tested on three separate occasions at three separate doses (placebo, 2.5 mg; 3.75 mg) and was categorized as a failure with the score of less than sixteen on both erotic and neutral video segments. He experienced yawning with both 2.5 mg and the 3.75 mg and was successful with this trial for two months until he inadvertently increased the dose. Adverse effects occurred within five minutes post-dosing (nausea and vomiting, dizziness, double and blurred vision, diaphoresis, and ashen coloring). The patient refused to retry medication after this incident. He stated he did not like this formulation.

Patient No. 2 was twenty-one year old male with erectile problems of a duration of three years. He had failed a previous course of yohimbine HCl. Ten minutes post-dosing with apomorphine at 2.5 mg he experienced yawing for a total of five yawns, and then experienced immediately major hemodynamic adverse effects. These included pale and ashen coloring, diaphoresis, nausea and vomiting, blurred vision, hypotension with a blood pressure of 70/50. Twenty minutes post adverse effect, vital signs were stable. The patient was feeling well, and coloring was good. This patient was then dropped from further testing.

Although the intranasal administration was effective in eliciting an erection, further testing of this intranasal formulation of apomorphine was discontinued because of possible overdose and increased side effects. The foregoing experience illustrates the need for reliable and relatively safer dosage forms.

Pilot Study #4

New sublingual tablet formulations of apomorphine at 3, 4 and 5 mg doses (Table I, above) were evaluated. Patients were tested on at least three separate occasions on at least three separate doses (placebo; 3 mg; and 4 mg). A 5 mg sublingual dose was also tested in some patients. The results of this study are summarized in Tables VIII and IX A-C, below.

To date, twelve patients have been completely evaluated on this formulation. All patients reported erectile dysfunction for more than two years. The patients' age range was thirty-nine to sixty-six years. Three patients had been successful with yohimbine HCl in the past, and two had previously not tried this compound. Seven patients of this group of twelve had previously failed a trial of yohimbine HCl. Of this latter group of seven, four were successfully treated with apomorphine.

Eight (67%) have been successful with apomorphine to date. Four (33%) were failures with apomorphine. Both 3 mg and 4 mg doses produced erectile responses. Several patients went on to a trial of the 5 mg sublingual dose which did not appear to be more effective than the relatively lesser doses in terms of erectile response. All eight of the successful patients continued on with the domestic use for a time period of one to four months. All patients reported good erectile activity and no side effects.

Analysis of visual analogue scales, both pre- and post-dosing with apomorphine, again indicated that the patients were relaxed but not sedated, and did not have feelings of arousal or anxiousness post-dosing.

5 The new formulations tested (3 mg; 4 mg; and 5 mg) were devoid of adverse effects. The patients felt well post testing, and did not report or demonstrate any adverse effects that had traditionally been seen with the administration of previous apomorphine liquid and  
10 intranasal preparations (Pilot Studies No. 1 and No. 3). The primary effect of yawning was still reported and observed at all doses, but the number and frequency of yawns was small (one or two).

The foregoing pilot study shows that 3-mg,  
15 4-mg and 5-mg apomorphine doses are effective in inducing penile erections, and also that there are no serious adverse effects with these preparations. Domestic use of these preparations was well accepted by patients and their partners. They were content with the  
20 convenience of dosing approximately fifteen minutes prior to sexual activity. All patients have stated that this was more acceptable than dealing with dosing on a routine basis.

**TABLE VI**  
**Inclusion/Exclusion Criteria**

**INCLUSION CRITERIA:**

- 5
1. Age 18-66 years.
  2. NPT circumference increase of 1.5 cm or more on one night and >70% rigidity.
  3. ICI circumference increase of 1.5 cm or more and >70% rigidity.

**EXCLUSION CRITERIA:**

- 10
1. Currently severe or life threatening systemic disease.
  2. Clinically significant ECG abnormalities.
  3. Personal or first degree family history of epilepsy.
  4. Abnormal:               Hepatic/renal function  
                                  Hematology
  5. Low:                    pre-trial testosterone  
   Low or High:        LH  
   High:                 Prolactin
  6. Hypertension requiring treatment.
  7. History of depression requiring treatment with antidepressants, ECT, or hospitalization.
  - 15 8. Symptomatic ischemic heart disease/or MI within the last three months.
  9. Diabetes.
  10. Failure to obtain informed consent.
  11. Legal cases.
  12. Unable or unwilling to comply with protocol.
  - 20 13. Drinks more than (on average) 45 units alcohol per week/or uses illicit drugs.
  14. History of syncope.
  15. Prohibited Drugs: sympathetic or parasympathetic types drugs, Beta blockers, Vasodilators, psychotropic medications, tranquilizers, thiazides, Captopril, Verapamil, Furosemide, Spironolactone, Metoclopramide, Cimetidine or other drugs which are likely to influence erectile function.



**TABLE VII**  
**Response to Erotic Videotape**

**1. Maximum increase in penile circumference**

5	<u>Circumference (cms.)</u>	<u>Score</u>
	0 - <0.5 cm.	0
	0.5 - <1.0 cm.	1
	1.0 - <1.5 cm.	2
	1.5 - <2.0 cm.	3
10	2.0 - <2.5 cm. lasts <1 min.	4
	2.5 or more lasts <1 min.	5
	2.0 - <2.5 cm. lasts at least 1 min.	6
	2.5 or more lasts at least 1 min.	7
	3.0 or more lasts at least 5 min.	8
15	3.0 or more lasts at least 10 min.	9

Score

A. Maximum increase in penile tip circumference \_\_\_\_\_

B. Maximum increase in penile basal circumference \_\_\_\_\_

**2. Maximum penile rigidity**

20	<u>Rigidity (%)</u>	<u>Score</u>
	0 - <10	0
	10 - <20	1
	20 - <30	2
	30 - <40	3
25	40 - <50	4
	50 - <60	5
	60 - <70	6
	70 - <80	7
	80 - <90	8
30	90 - 100	9

Score

C. Maximum penile tip rigidity \_\_\_\_\_

D. Maximum penile basal rigidity \_\_\_\_\_

**3. Total score (A, B, C & D)**

35 **A score of less than 16 indicates erectile dysfunction**

TABLE VIII  
Summary of Results from Pilot Study #4 in Psychogenic Patients

Patient # (Wt., kg)	PLACEBO		Apomorphine • HCl Sublingual Tablet							
	Erotic #1	Neutral #1	3 Mg Dose (µg/kg)		4 Mg Dose (µg/kg)		5 Mg Dose (µg/kg)		Erotic #4	Neutral #4
			Erotic #2	Neutral #2	Erotic #3	Neutral #3	Erotic #3	Neutral #3		
401 (68.5)	31	28	29 (44)	27 (44)	33 (58)	27 (58)				
402 (70.3)	12	4	12 (43)	4 (43)	17 (57)	6 (57)				
403 (118)	16	4	22* (25)	5 (25)	22* (34)	25 (34)				
404 (83.5)	24	10	26* (36)	17 (36)	25* (48)	17 (48)				
405 (78)	11	1	18* (38)	6 (38)	12 (51)	8 (51)			10 (64)	5 (64)
406 (80)	14	5	18* (38)	17 (38)	17* (50)	2 (50)				
407 (100)	8	0	18* (30)	4 (30)	10 (40)	3 (40)				
408 (86.2)	28	18	32 (35)	21 (35)	34 (46)	22 (46)				
409 (93)	2	0	4 (32)	1 (32)	8 (43)	6 (43)			5 (54)	4 (54)
410 (80)	3	0	13 (38)	16 (38)	8 (50)	7 (50)				
411 (98)	13	5	26* (31)	23* (31)	24* (42)	20 (42)				
412 (73)	7	3	7 (41)	1 (41)	28* (55)	19* (55)				

\* Patients with score higher than 16 (see scoring table) are positive respondents.

Out of 12 patients who were treated in this study, 5 showed improvement at both 3 mg and 4 mg doses. Two (2) showed response only at one dose.

No improvement in clinical response was observed at 5 mg dose.

The data of Pilot Study #4 were analyzed in two ways. First, mean erectile function was compared across placebo, 3 mg and 4 mg doses under two stimulus backgrounds, erotic and neutral. Next erectile function scores were dichotomized, with values less than sixteen considered to reflect erectile insufficiency.

A. Mean Erectile Function

Table IX A shows means and standard errors for all three treatments under both backgrounds, erotic and neutral. Means were compared using a restricted maximum likelihood generalized linear model containing two main effects, treatment and stimulus, and the treatment by stimulus interaction. An appropriate variance-covariance structure was established for the underlying statistical model using Akaike's criterion. Table IX B presents the statistical results for the main effects of treatment and of stimulus, for the treatment by stimulus interaction, and for orthogonal contrasts within the erotic and neutral conditions. It can be seen that the treatment main effect (i.e., general difference across treatment conditions without regard to stimulus background) is statistically significant; that the main effect of stimulus (i.e., general difference across stimulus backgrounds without regard to treatment) is statistically significant; and that the treatment by stimulus interaction is not statistically significant. These findings imply that active treatment is more effective than placebo and that this finding, although stronger when using an erotic stimulus, is true regardless of stimulus background (see FIGURE 1). The orthogonal (statistically independent) contrasts confirm that active treatment is superior at a statistically significant level under both erotic and neutral conditions, but also indicate that the difference

between the 3 mg and 4 mg dose does not exceed that expected by chance for the number of patients (12) used in this study.

B. Percent Successful Erectile Function

FIGURE 2 and Table IX C show that the statistically significant superiority of active over placebo treatment, regardless of stimulus background, is maintained when the erectile function scores are classified to reflect success (score at least 16) or failure (score less than 16).

**TABLE IX A**  
**Mean and Percent Successful Erectile Function**

Stimulus	Treatment	N	Mean (SE)	Percent (SE)
Erotic	Placebo	12	14.08 (2.69)	33.33 (13.61)
	3 mg	12	18.75 (2.51)	66.67 (13.61)
	4 mg	12	19.83 (2.67)	66.67 (13.61)
Neutral	Placebo	12	6.50 (2.45)	16.67 (10.76)
	3 mg	12	11.83 (2.68)	50.00 (14.43)
	4 mg	12	13.50 (2.61)	50.00 (14.43)

Note: Mean (SE) from SAS PROC UNIVARIATE. Percent (SE) from SAS PROC CATMOD.

**TABLE IX B**  
**Anova for Mean Erectile Function**

EFFECT	DF	F	P-value
Treatment	2.66	11.56	0.0000
Stimulus	1.66	37.14	0.0000
Treatment by Stimulus	2.66	0.10	0.9046
Contrasts			
Erotic: Placebo vs. Treatment	1.66	9.30	0.0033
Erotic: 3 mg vs. 4 mg	1.66	0.30	0.5849
Neutral: Placebo vs. Treatment	1.66	13.03	0.0006
Neutral: 3 mg vs. 4 mg	1.66	0.71	0.4014

Note: Restricted maximum likelihood analysis performed using SAS PROC MIXED.

**TABLE IX C**  
**Logistic Regression for Percent**  
**Successful Erectile Function**

EFFECT	DF	X <sup>2</sup>	P-value
Treatment	2	15.36	0.0005
Stimulus	1	5.14	0.0233
Treatment by Stimulus	2	0.00	1.0000
Contrasts			
Erotic: Placebo vs. Treatment	1	9.60	0.0019
Erotic: 3 mg vs. 4 mg	1	0.00	1.0000
Neutral: Placebo vs. Treatment	1	9.60	0.0019
Neutral: 3 mg vs. 4 mg	1	0.00	1.0000

Note: Analysis performed using SAS PROC CATMOD.

**TABLE X**  
**Visual Analogue Scale (VAS)**  
(to be completed by the patient)

Please mark each line clearly at the point which indicates how you are feeling right now.  
Each line represents the full range of each feeling. (There are no right or wrong answers)

				Score (mm)
1.	Alert	_____	Drowsy	_____
2.	Calm	_____	Excited	_____
3.	Yawning	_____	Not Yawning	_____
4.	Fuzzy	_____	Clear Headed	_____
5.	Well Coordinated	_____	Clumsy	_____
6.	Tired	_____	Energetic	_____
7.	Contented	_____	Disconnected	_____
8.	Troubled	_____	Tranquil	_____
9.	Mentally slow	_____	Quick Witted	_____
10.	Tense	_____	Relaxed	_____
11.	Attentive	_____	Dreamy	_____
12.	Stomach Upset	_____	Feeling Well	_____
13.	Anxious	_____	Carefree	_____

(measure from left to right)

### Dose Evaluation Study

Clinical response to sublingual administration of apomorphine was evaluated utilizing a group of 60 non-vasculogenic impotent patients. Each patient had a history of erectile dysfunction for at least 3 months, normal biothesiometry response, and normal cavernosometry results.

The patients were divided into seven groups. Each group received a predetermined dosage of apomorphine for 20 days in the form of apomorphine hydrochloride tablets 20 minutes prior to intercourse. Seven different dosages were evaluated - 3 mg, 4 mg, 5 mg, 6 mg, 7 mg, 8 mg and 10 mg. The tablet constituents were those shown in Table I, above. Assessment of response was made on the basis of the patient's report of his experience. A response was deemed positive when the patient experienced an erection sufficiently rigid to effect penetration. Side effects such as nausea and/or vomiting, if present, were noted as well.

The results of this study are compiled in Table XI, below.

TABLE XI  
Results of Dose Evaluation Study

No. of Patients	Dosage, mg	Positive Responses		Nausea		Vomiting	
		No.	%	No.	%	No.	%
5	3	0	0	0	0	0	0
5	4	2	40	1	20	1	20
10	5	5	50	2	20	1	10
10	6	7	70	2	20	2	20
10	7	7	70	2	20	2	20
10	8	7	70	3	30	3	30
10	10	8	80	4	40	4	40

From the foregoing Table it can be seen that at a 4-mg dosage 40 percent of patients had a positive response, at a 5-mg dosage 50 percent of patients had a positive response, at 6-mg, 7-mg, and 8-mg dosages 70 percent of patients had a positive response and at a 10-mg dosage 80 percent of patients had a positive response. However, the incidence of side effects increased as well as the dosage was increased.

The aforesaid apomorphine dosage forms are also well suited for diagnosing male human patients suffering from male erectile dysfunction. For diagnostic purposes, at least about 3 milligrams of apomorphine are administered sublingually to the patient and the patient is exposed to a visual erotic stimulus, e.g., an erotic videotape, while the patient's response thereto is monitored. If deemed desirable for diagnostic purposes, up to about 10 milligrams of apomorphine can be administered to the patient.

In particular, the patient's maximum increase in penile circumference (preferably tip as well as basal) is determined and the patient's maximum penile rigidity (preferably tip as well as basal) is determined. The determined circumferential increase and rigidity values are then compared against a predetermined base value. Equivalent methods of determining tumescence and rigidity can also be utilized.

#### Pilot Study #5

A clinical study, "Absorption and pharmacokinetic evaluation of apomorphine after sublingual and intravenous routes of administration" compared the absorption and pharmacokinetic profile of apomorphine administered intravenously and slowly at a 1 mg dose with apomorphine sublingual tablets at doses of

4 mg (Table I) and 8 mg administered on 3 occasions, 4 days apart, over a 12 day period in a cross-over study design. The tolerance for apomorphine for each route and each dose administered was determined.

5           The study was conducted as an open-label, single center, 3-way crossover design. The study population was seven healthy, Caucasian male volunteers between 18 and 35 years of age. A 15-day pre-study evaluation period was followed by a 12-day active  
10 treatment phase. Three doses (one intravenous; 2 sublingual) were administered to each subject in random order 4 days apart. A total of 36 serum samples were obtained from each subject at the following time  
15 periods: 0, 2, 3, 5, 10, 20, 30 and 45 minutes; and 1, 2, 3, 4 and 6 hours post dose administration.

Safety was assessed within 15 days prior to study start and within one week after the last dose was administered. General physical examination was performed. Change from baseline in vital signs,  
20 height/weight measurements, ECG, orthostatic arterial pressures, heart rate, serum chemistry profile, hematology profile and urinalysis were recorded. Adverse experiences were recorded at each visit and tabulated.

### **Data Analysis**

#### **25   1.   Pharmacokinetic Analysis**

Pharmacokinetic analysis was performed by compartmental and noncompartmental methods described below. Nonlinear, iterative, least-squares regression analysis was performed with the computer program, PPHARM  
30 (Simed Co., Philadelphia, PA).



### A. Compartmental Analysis

The apomorphine plasma concentration data for each subject following intravenous administration was fitted to two-compartment open model with a first order input function as described by the following equations.

Plasma apomorphine concentration was described for intravenous administration data by equation (1):

$$C_t = Ae^{-\alpha t} + Be^{-\beta t} \quad (1)$$

Plasma apomorphine concentration was described for sublingual tablet administration by equations (2) and (3):

$$C_t = \frac{FDk_a}{V_d(k_a - k_e)} (e^{-k_e(t-t_{lag})} - e^{-k_a(t-t_{lag})}) \quad (2)$$

$$C_t = Ae^{-\alpha(t-t_{lag})} + Be^{-\beta(t-t_{lag})} + Ce^{-k_a(t-t_{lag})} \quad (3)$$

In the above equations,  $C_t$  is the apomorphine plasma concentration at time  $t$ ;  $F$  is the relative bioavailability, which is assumed to be one for intravenous administration;  $K_a$  is the first order rate constant for sublingual absorption;  $K_e$  is the first order rate constant for elimination;  $V_d$  is the volume of distribution;  $D$  is the apomorphine dose;  $t$  is time;  $t_{lag}$  is the lag time before onset of sublingual absorption;  $A$ ,  $B$ ,  $C$  are the intercepts of the distribution, elimination, and absorption phases, respectively;  $\alpha$  is the distribution rate constant;  $\beta$  is the elimination rate constant; and  $K_a$  is the absorption rate constant.

Initial estimates of the intravenous and sublingual pharmacokinetic parameters were obtained with the computer program PPHARM (Simed Co., Philadelphia, PA). These initial estimates were used to fit the data to equations (1), (2) and (3) by nonlinear iterative least squares regression analysis. The results are shown graphically in FIGURE 4. The estimate of F (relative bioavailability) for sublingual administration was obtained from the noncompartmental analysis outlined below.

Visual inspection of the fitted curves, analysis of the residual plots, the Akaike information criterion, and correlation coefficients between observed and calculated values were used to select the appropriate pharmacokinetic model for each set of plasma concentration versus time data. A weighting factor was used to fit the data.

The regression analysis provided the final estimates of the pharmacokinetic parameters:  $V_d$ ,  $K_e$ ,  $K_a$ , and  $t_{lag}$  from equation (2), and A, B, C,  $\alpha$ ,  $\beta$ ,  $k_a$ , and  $t_{lag}$  from equation (3). The maximum plasma concentration ( $C_{max}$ ), time to maximum plasma concentration ( $T_{max}$ ), and  $V_d$  (volume of distribution) were calculated using standard compartmental pharmacokinetic equations (Gibaldi, M. & Perrier, D. Pharmacokinetics, 2d edition, Marcel Dekker, Inc. New York, 1982).

The values for  $C_{max}$  and  $T_{max}$  obtained by visual inspection of the plasma concentration versus time curve were reported for comparative purposes. Model dependent and independent pharmacokinetic parameters (presented in Tables XII-XV) were calculated for each patient using the best fit of Equation (3) to the data.

## B. Noncompartmental Analysis:

The area under the curve,  $AUC_{0-inf}$ , was determined by adding the  $AUC_{0-last}$  to the  $AUC_{t-last}$ , where  $AUC_{t-last} = C_t / R_e$ ,  $C_t$  is the plasma concentration at time  $t$ , and  $R_e$  is  $K_e$ , the first order rate constant for elimination.

If the plasma concentration versus time data for a subject could not be adequately fit to equation (1), (2) or (3), the  $K_e$  was determined by linear regression analysis of the log plasma concentration versus time during the post-absorption phase. Estimates of noncompartmental parameters  $C_{max}$  and  $T_{max}$  were obtained from visual inspection of the plasma concentration time curves.

The relative bioavailability (F) for a sublingual dose was calculated by the following equation:

$$F = \frac{AUC_{SL} * D_{IV}}{AUC_{IV} * D_{SL}} \quad (4)$$

## 2. Statistical Analysis:

An analysis of variance (ANOVA) for a three-way crossover study design was utilized to compare the pharmacokinetic parameters ( $AUC$ ,  $C_{max}$ , and  $t_{lag}$ ) determined as described above for the intravenous administration at 1 mg dose and those determined for the sublingual administration of apomorphine at the 4 mg and the 8 mg doses. The ANOVA was tested for the presence of any period or residual carryover effects in the data. Statistical significance was set at an alpha level equal to 0.05. The ability of the ANOVA to detect both a 20% difference and the observed difference between the

5 sublingual and intravenous pharmacokinetic parameters was determined. In addition to the ANOVA, the 95% confidence interval of the percent difference between the sublingual and SC parameters was calculated from the error variance and degrees of freedom of the ANOVA model.

The data were summarized as the mean  $\pm$  standard deviation in Tables XII-XV below.

1004453 10304 885400

TABLE XII  
NONCOMPARTMENTAL PHARMACOLOGIC PARAMETERS  
(MEAN ± SD) AND RANGE

Parameter	IV Administration				4 mg Dose				8 mg Dose			
	Mean	±SD	Low	High	Mean	±SD	Low	High	Mean	±SD	Low	High
Ke (min <sup>-1</sup> )	0.0237	0.0140	0.0091	0.0432	0.0156	0.0138	0.0038	0.0336	0.0056	0.0036	0.00222	0.0102
T <sub>1/2</sub> (min)	39.44	0.219	16.04	76.49	89.18	75.43	20.62	183.60	176.30	112.30	68.09	314.6
Tmax (min)	2.286	1.254	1.000	5.000	17.50	18.48	5.000	45.00	52.50	85.10	5.000	15.00
Cmax (min)	8.364	3.886	3.400	12.90	0.8375	0.6848	0.3000	0.8500	2.069	2.366	0.5750	1.150
AUC (0-inf) (min*ng/ml)	206.9	45.47	140.8	260.1	31.64	18.62	10.13	55.55	339.9	459.2	15.00	316.6
Cl (ml/min)	0.0051	0.0012	0.0038	0.0071	0.0456	0.0361	0.0180	0.0988	0.2056	0.2460	0.0253	0.5333
Vd (beta) (ml)	0.2344	0.1532	0.0056	0.4982	4.076	2.053	1.017	5.277	69.09	115.80	7.932	64.27
Vd (SS) (ml)	0.1942	0.0817	0.1357	0.3401	1.836	0.7112	0.999	2.475	46.30	74.46	6.523	12.42
MRT (min)	40.29	18.90	20.14	75.32	64.25	55.14	15.12	137.5	143.7	148.0	23.29	329.3
F' (% Relative Bioavailability)	-	-	-	-	0.04	(4.0%)	-	-	0.21	(21%)	-	-
F' = (AUC <sub>SL</sub> *DOSE <sub>IV</sub> )/(AUC <sub>IV</sub> *DOSE <sub>SL</sub> )												

Cl = clearance  
Vd = volume of distribution @ 0.5 stage  
Vd (SS) = volume of distribution steady state  
MRT = means residual time

**TABLE XIII**  
**Noncompartmental Pharmacokinetic Parameters (Mean  $\pm$  SD) for IV Administration (1 mg)**  
**n=7**

	Subject				Subject			Range			
	#1	#2	#3	#4	#5	#6	#7	Mean	$\pm$ SD	Low	High
Ke (/min)	0.0432	0.0129	0.0091	0.0419	0.0168	0.0150	0.0268	0.0237	0.0140	0.0091	0.0432
T <sub>1/2</sub> (min)	16.04	53.79	76.49	16.56	41.22	46.18	25.84	39.44	21.92	16.04	76.49
Tmax (min)	2.000	2.000	2.000	2.000	2.000	5.000	1.000	2.286	1.254	1.000	5.000
Cmax (ng/ml)	8.400	11.200	4.150	12.250	12.900	3.400	6.250	8.364	3.886	3.400	12.900
AUC (0-inf) (min*ng/ml)	140.8	255.4	221.5	177.1	224.6	169.0	260.1	206.9	45.47	140.8	260.1
Cl (ml/min)	0.0071	0.0039	0.0045	0.0056	0.0045	0.0059	0.0038	0.0051	0.0012	0.0038	0.0071
Vd (beta) (ml)	0.1643	0.3039	0.4982	0.0056	0.2648	0.2574	0.1466	0.2344	0.1532	0.0056	0.4982
Vd (SS) (ml)	0.1430	0.1512	0.3401	0.1147	0.2174	0.2574	0.1357	0.1942	0.0817	0.1357	0.3401
MRT (min)	20.14	38.62	75.32	20.32	48.82	43.50	35.31	40.29	18.90	20.14	75.32

**TABLE XIV**  
**Noncompartmental Pharmacokinetic Parameters (Mean  $\pm$  SD)**  
**for Sublingual Administration (4mg Dose)**

n=4

	Subjects					Range			
	#1	#3	#6	#7	Mean	$\pm$ SD	Low	High	
Ke (/min)	0.0188	0.0336	0.0060	0.0038	0.0156	0.0138	0.0038	0.0336	
T <sub>1/2</sub> (min)	36.78	20.62	115.7	183.6	89.18	75.43	20.62	183.6	
Tmax (min)	10.00	10.00	5.000	45.00	17.50	18.48	5.000	45.00	
Cmax (ng/ml)	0.8500	1.8000	0.3000	0.4000	0.8375	0.6848	0.3000	0.8500	
AUC (0-inf) (min*ng/ml)	10.13	29.25	31.64	55.55	31.64	18.62	10.13	55.55	
Cl (ml/min)	0.0988	0.0342	0.0316	0.0180	0.0456	0.0361	0.0180	0.0988	
Vd (beta) (ml)	5.241	1.017	5.277	4.769	4.076	2.053	1.017	5.277	
Vd (SS) (ml)	1.494	0.999	2.377	2.475	1.836	0.7112	0.999	2.475	
MRT (min)	15.12	29.23	75.19	137.47	64.25	55.14	15.12	137.47	

**TABLE XV**  
**Noncompartmental Pharmacokinetic Parameters (Mean  $\pm$  SD)**  
**for Sublingual Administration (8mg Dose)**

n=4

	Subjects						Range			
	#2	#3	#4	#6	Mean	$\pm$ SD	Low	High		
Ke (/min)	0.0067	0.0032	0.0120	0.0022	0.0056	0.0036	0.0022	0.0102		
T <sub>1/2</sub> (min)	104.0	218.4	68.09	314.6	176.3	112.3	68.09	314.6		
Tmax (min)	180.0	10.00	15.00	5.000	52.50	85.10	5.000	180.0		
Cmax (ng/ml)	5.600	1.150	0.9500	0.5750	2.069	2.366	0.5750	5.600		
AUC (0-inf) (min*ng/ml)	996.6	316.6	31.25	15.00	339.9	459.2	15.00	996.6		
Cl (ml/min)	0.008	0.0253	0.2560	0.5333	0.2056	0.2460	0.008	0.5333		
Vd (beta) (ml)	1.204	7.932	25.15	242.1	69.09	115.8	1.204	242.1		
Vd (SS) (ml)	157.9	8.320	6.523	12.42	46.30	74.46	6.523	157.9		
MRT (min)	196.7	329.3	25.48	23.29	143.7	148.0	23.29	329.3		



Summary of Pharmacokinetic Parameters for Apomorphine HCl in Humans

THIS STUDY										PUBLISHED DATA								
										Durif, F. et al., Clin. Neuropharm. 16:157-166 (1993)			Gancher, S.T., et al. <i>Movement Disorders</i> 6:212-216 (1991).			Montastruc, J.L., et al. <i>Clin.Neuropharmacol.</i> 14:432-437 (1991).		
										s.l.	s.l.	s.l.	s.l.	s.c.	i.v.	s.l.	s.c.	
ROUTE	i.v.	s.l.	s.l.	s.l.	s.l.	s.l.	s.l.	s.c. <td>i.v.<td>s.l.</td><td>s.c.<td>s.l.</td><td>s.c.</td></td></td>	i.v. <td>s.l.</td> <td>s.c.<td>s.l.</td><td>s.c.</td></td>	s.l.	s.c. <td>s.l.</td> <td>s.c.</td>	s.l.	s.c.					
# Subjects	7	7	7	7	7	7	7	5	5	9	9	9	9					
# Tablets x Strength (mg)	n/d	1 x 4	1 x 8	1 x 8	14 x 3	7 x 3	14 x 3	3 x 6	n/a	n/a	10 x 3	n/a	n/a					
Dose (mg/kg)	0.01	0.06	0.114	0.3	0.6	0.25	0.02	0.038	0.42	0.04								
C <sub>max</sub> (ng/ml)	8.3	0.83	2.07	7.5	22.7	14.3	19.36	31.2	28	26								
T <sub>max</sub> (min)	2.2	17.5	52.5	31.5	38.3	45	6.5	6.7	41	18								
AUC (min*ng/ml)	207	31.6	340	929	2,277	1,057	592.7	881.1	1,882	837								
Cl (l/hr/kg)	4.37	n/d	n/d	2.1	1.8	n/d	n/d	n/d	n/d	n/d								
Vd (l/kg)	3.35	2.33	2.07	3.4	2.8	n/d	n/d	0.043	n/d	n/d								
MRT (min)	40.3	64.2	143.7	128	125	n/d	n/d	n/d	n/d	n/d								
T <sub>1/2</sub> (min)	39.4	89.2	176.3	72	70	n/d	n/d	n/d	n/d	n/d								
Bioavailability (F)	n/a	4%	21%	10%	10%	17%	n/a	n/a	n/d	n/a								

n/d = not done

n/a = not applicable

\*Calculated

Clin. Neuropharm. 16:157-166 (1993)

The results, summarized in Tables XII-XV above and shown graphically in FIGURE 4, show that the plasma concentration of apomorphine drops quickly when the drug is administered intravenously. In contrast, the plasma concentration of apomorphine rises slowly to a lower level when administered sublingually.

The importance of these findings is put into perspective when compared to information on the administration of apomorphine that is available in the literature (Table XVI). The sublingual administration of apomorphine by the present invention produced a lower plasma concentration than the administration and dosage regimes listed for previous reports.

#### Pilot Study #6

A clinical study of patient tolerance of escalating doses in sublingual tablet administration of APO for the treatment of psychogenic male erectile dysfunction was performed. The pilot study compared the effects of sublingual tablet administration of placebo, and 4, 6 and 8 mg apomorphine hydrochloride (APO) on male erectile dysfunction as measured by RIGISCAN™ monitoring and self-reported satisfaction with the treatment results.

The study included 50 men with psychogenic male erectile dysfunction (MED). The study was conducted in three phases. In the first phase, the subject's penile erectile response (measured with the RIGISCAN™ ambulatory tumescence monitor) was evaluated. The subject received a placebo tablet for sublingual administration and then viewed a 30-minute video consisting of two 10-minute erotic sequences separated by a 10-minute neutral sequence. Subjects completed a visual analogue scale questionnaire (VAS, Table VII) about their feelings and well-being.

In the second phase, subjects returned to the clinic for four visits, each visit one week apart. Subjects received one sublingual administration of either placebo, or 4, 6 and 8 mg APO at each visit. Doses of APO were administered in ascending order with the placebo being randomly assigned for use at one of the four visits. The procedures performed before and after drug administration were the same as those in the first phase. After completion of the fourth visit, the investigator determined for each subject the most effective and well-tolerated APO dose for home use in the third phase of the study.

The third phase, a home-use phase, lasted 5 weeks. During this phase, subjects attempted coitus at least once each week after taking a single APO tablet. After each attempt the subject and his partner completed a Sexual Function questionnaire (Table XVII). Subjects had a final evaluation at the end of the 5-week, home-use phase.

Fifty males with psychogenic MED were enrolled in this three phase trial. The first aim of this study was to determine the safety and tolerance of APO in the treatment of MED. Several adverse events directly linked with administration of APO in humans were expected: yawning, nausea, vomiting, and cardiovascular effects. Indeed, nausea was the primary adverse event reported in this trial with an overall incidence of less than 13% of the subjects for all administered doses and only two cases were considered severe. The incidence of vomiting was less than 3% for all administered doses.

Hypotension was reported as an adverse event in some subjects in this study, along with bradycardia, dizziness, syncope, and pallor. Only single cases of hypotension and pallor were judged severe in this study.

Increased sweating and fatigue were also reported. One of the cases of increased sweating was considered severe. The other severe adverse events (mouth edema, dysphagia, upper respiratory tract infection) were judged unrelated to treatment.

Changes in the serum chemistry values and vital signs paralleled the adverse event reports. There were no clinically significant changes except for one subject judged to have abnormal liver function of unknown origin. There were no clinically significant changes in the blood or urinalysis values due to drug.

The efficacy of APO was evaluated during the first two phases of the study in which subjects were attached to the RIGISCAN<sup>TM</sup> monitor. Subjects were initially treated with placebo in the first phase. In the second phase, patients received 4, 6 and 8 mg APO tablets with a placebo tablet randomly interspersed in the treatment.

There were highly significant effects of APO treatment compared to placebo. These observations indicate that APO has effects on penile function in both erotic and neutral environments (Tables XVIII-XX). All summed scores showed significant treatment effects at one or more of the three doses of APO. The overall RIGISCAN<sup>TM</sup> score results were significant to highly significant for a treatment effect of 4, 6 and 8 mg compared to the initial placebo. In addition, most of the treatment effects were significant to highly significant compared to the second placebo.

The effects in the erotic video sequences were larger than the effects in the neutral video sequence (TABLES XVIII-XX). Effects in erotic video sequence one were larger than the effects in erotic video sequence two (TABLE XVIII). More significant treatment effects

were seen in response to the neutral video sequence, but this reflects the larger number of subjects in this data subset, as one center did not show the erotic video sequences. All doses of APO were effective in causing  
5 erections (RIGISCAN™ reading  $\geq 15$  in the presence of erotic stimulation; TABLE XVIII).

During the third phase, subjects had recorded at baseline, their satisfaction, erection, number of attempts, and successful intercourse on a VAS scale.  
10 Evaluable subjects first recorded a success rate, then completed VAS for erection results and satisfaction with intercourse following take-home treatment. Success rate was calculated for mg as well as  $\mu\text{g/kg}$  body weight doses (males). Several evaluations of the data were made  
15 including the male and female responses to treatments. The overall average success rate is 69% with APO treatment which is much higher than the average baseline rate 28% (Tables XXI, XXII).

The success rate showed numerical increase at  
20 tablet strength from 4 mg to 6 mg, but a decrease at 8 mg (TABLE XXI). The highest success rate was 73% in both males and females at a tablet strength of 6 mg (TABLE XXI). When the dosage is examined as a function of body weight, a dosage range of 50-74  $\mu\text{g/kg}$  gave the  
25 highest success rate (: 82%) in females and (80%) in males (Table XXII). The dosage range of 35-50  $\mu\text{g/kg}$  gave the highest success rate.

The optimal response was observed with 4 or 6  
30 mg APO sublingual tablets which caused erections in the majority (72%) of men with male erectile dysfunction (MED) with few severe adverse effects.

**TABLE XVII**  
**SEXUAL FUNCTION STUDY HOME QUESTIONNAIRE - Male**  
Please answer questions within 12-24 hours of taking sublingual tablet.

5

Initials: \_\_\_\_\_ Subject#: \_\_\_\_\_

Today's Date: \_\_\_\_\_ Time: \_\_\_\_\_

Date Tablet Taken: \_\_\_\_\_ Time: \_\_\_\_\_

10

The lines below represent the full range of feeling or response. Please mark each line clearly with a vertical (straight up and down) stroke at the point which represents your response. (There are no right or wrong answers. Do not write in boxes on right.)

15

1. What was your erection result after taking the sublingual tablet?

No  
Erection \_\_\_\_\_

Rigid Erection  
Suitable for  
Penetration [ ]

20

2. Did you have intercourse with [ ] Yes [ ] No  
wife/partner after taking tablet?

IF NO, please circle  
all reasons that apply:

0 - No erection.

1 - Erection not sufficient for penetration.

2 - Felt sick after taking tablet. (Describe below in #4.)

3 - I decided not to participate in intercourse.

4 - Wife/partner decided not to participate.

5 - Unrelated interruption (example, telephone call).

6 - Wife/partner menstruating.

7 - Other, explain: \_\_\_\_\_

25

30

3. What was your level of satisfaction with this attempt at sexual intercourse?

Extremely  
Unsatisfied \_\_\_\_\_

Extremely  
Satisfied [ ]

35

40

4. Please describe any adverse reactions you experienced after taking the sublingual tablet. (Indicate when the reaction started and stopped, and any intervention taken i.e. "nosebleed on 5/1/94, used a cold compress".)

45

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5. Other comments?

50

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TABLE XVIII  
Total RIGISCAN™ Scores by Phase

Mean ±SEM

Phase II

	Phase I		Phase II		
	Placebo 1	Placebo 2	4 mg	6 mg	8 mg
Video					
Erotic 1 N = 31-36	11.44 ± 1.77	13.38 ± 2.05	15.31 ± 1.76*	17.09 ± 1.64**	19.84 ± 1.61**
Erotic 2 N = 29-36	11.39 ± 1.70	13.31 ± 1.88	15.26 ± 1.72*	16.44 ± 1.98*	17.79 ± 1.96**
Neutral N = 41-48	7.98 ± 1.24	7.49 ± 1.26	11.11 ± 1.30**	12.76 ± 1.12**	11.98 ± 1.37**
Corresponding p-values (placebo 1/placebo 2)					
Erotic 1	--	0.3274	0.0120	0.0007	0.0001
	--	--	0.1405	0.0166	0.0005
Erotic 2	--	0.4013	0.0276	0.0196	0.0007
	--	--	0.1907	0.1365	0.0091
Neutral	--	0.6243	0.0230	0.0009	0.0060
	--	--	0.0074	0.0002	0.0017

\* Significantly higher than placebo 1

\*\* Significantly higher than placebo 1 and placebo 2

TABLE XIX

Penile Measurements (Maximum Increases Measured by RIGISCAN™), Erotic Video Sequence #1  
Repeated Measures Analysis of Variance

Site	Treatment Source	DESCRIPTIVE STATISTICS			ADJUSTED (LS) MEAN		-----ANALYSIS OF VARIANCE-----	
		N	MEAN	SEM	LSMEAN	SEM		
ALL SITES	Placebo #1	36	11.44	1.770	12.22	1.666	Treatment	0.0001 *
	Placebo #2	32	13.38	2.051	13.65	1.714	Site	0.0264 *
	4 mg	35	15.31	1.761	15.80	1.674	Treatment by Site	0.0595
	6 mg	34	17.09	1.841	17.20	1.695	4mg vs Placebo #1	0.0120 *
SITE #1	8 mg	31	19.84	1.610	19.11	1.745	6mg vs Placebo #1	0.0007 *
	ALL TREATMENTS	11	10.76	2.372	11.04	2.498	8mg vs Placebo #1	0.0001 *
	Placebo #1	11	9.73	2.854	9.73	2.931	4mg vs Placebo #2	0.1504
	Placebo #2	10	9.00	3.300	9.21	2.996	6mg vs Placebo #2	0.0166 *
SITE #2	4 mg	11	8.09	2.410	8.09	2.931	8mg vs Placebo #2	0.0005 *
	6 mg	11	10.82	3.065	10.82	2.931	Placebo #1 vs. #2	0.3274
	8 mg	9	17.89	2.988	17.36	3.070		
	ALL TREATMENTS	16	13.89	1.942	14.25	2.083		
SITE #4	Placebo #1	16	8.94	2.233	8.94	2.430		
	Placebo #2	14	11.71	2.768	11.38	2.515		
	4 mg	15	15.27	2.379	15.10	2.476		
	6 mg	15	17.60	2.267	17.43	2.476		
SITE #4	8 mg	15	18.60	2.265	18.43	2.476		
	ALL TREATMENTS	9	21.21	3.437	21.49	2.776		
	Placebo #1	9	18.00	4.304	18.00	3.240		
	Placebo #2	8	21.75	4.242	20.36	3.337		
SITE #4	4 mg	9	24.22	2.837	24.22	3.240		
	6 mg	8	24.75	3.740	23.36	3.337		
	8 mg	7	25.00	3.259	21.52	3.444		



TABLE XX

Penile Measurements (Maximum Increases Measured by RIGISCAN™), Neutral Video Sequence  
Repeated Measures Analysis of Variance

Site	Treatment Source	DESCRIPTIVE STATISTICS			ADJUSTED (LS) MEAN		-----ANALYSIS OF VARIANCE-----	
		N	MEAN	SEM	LSMEAN	SEM		
ALL SITES	Placebo #1	48	7.98	1.236	8.34	1.220	Treatment	0.0002
	Placebo #2	43	7.49	1.257	7.65	1.272	Site	0.1092
	4 mg	47	11.11	1.295	11.47	1.226	Treatment by Site	0.7176
	6 mg	45	12.76	1.116	13.10	1.268	4mg vs Placebo #1	0.0230
SITE #1	8 mg	41	11.98	1.366	12.40	1.331	6mg vs Placebo #1	0.0009
	ALL TREATMENTS	11	10.56	1.987	10.70	1.789	8mg vs Placebo #1	0.0060
	Placebo #1	11	8.91	2.470	8.91	2.494	4mg vs Placebo #2	0.0074
	Placebo #2	10	5.60	2.574	5.68	2.587	6mg vs Placebo #2	0.0002
SITE #2	4 mg	11	10.45	1.965	10.45	2.494	8mg vs Placebo #2	0.0017
	6 mg	11	12.73	2.832	12.73	2.494	Placebo #1 vs. #2	0.6243
	8 mg	9	16.22	3.099	15.73	2.692		
	ALL TREATMENTS	16	7.02	1.192	7.22	1.495		
SITE #3	Placebo #1	16	4.44	1.554	4.44	2.068		
	Placebo #2	14	5.86	2.099	5.71	2.182		
	4 mg	15	8.73	2.610	8.70	2.126		
	6 mg	15	9.60	1.514	9.56	2.126		
SITE #4	8 mg	15	7.73	1.694	7.70	2.126		
	ALL TREATMENTS	12	12.22	1.476	12.09	1.706		
	Placebo #1	12	11.33	2.244	11.33	2.388		
	Placebo #2	11	10.00	1.902	10.61	2.469		
SITE #4	4 mg	12	11.83	2.564	11.83	2.388		
	6 mg	12	13.58	1.794	13.58	2.388		
	8 mg	11	12.45	2.458	13.07	2.469		
	ALL TREATMENTS	9	11.63	2.864	12.35	2.023		
SITE #4	Placebo #1	9	8.67	4.052	8.67	2.758		
	Placebo #2	8	9.25	3.990	8.58	2.891		
	4 mg	9	14.89	3.071	14.89	2.758		
	6 mg	7	18.14	2.747	16.51	3.046		
SITE #4	8 mg	6	15.33	4.462	13.11	3.236		

TABLE XXI

Reported Success by Tablet Strength

Group	4 mg	6 mg	8 mg	Overall
Female	5/7 (71.4%)	11/15 (73.3%)	4/7 (57.1%)	20/29 (69.0%)
Male	5/7 (71.4%)	11/15 (73.3%)	4/7 (57.1%)	20/29 (69.0%)

TABLE XXII

Reported Success by Apomorphine Dosage ( $\mu\text{g/kg}$ )

Group	35-50 $\mu\text{g/kg}$	50-74 $\mu\text{g/kg}$	>74 $\mu\text{g/kg}$	Overall
Female	3/5 (60.0%)	9/11 (81.8%)	8/13 (61.5%)	20/29 (69.0%)
Male	4/5 (80.0%)	8/11 (72.7%)	8/13 (61.5%)	20/29 (69.0%)

Subject Evaluability Rules for Take-home Part

1. Subjects who get one out of two successful intercourse is considered a success [based on subject's answers to the take-home questionnaires].
2. Subjects who tried the study medication at home, for at least two times.
3. Subjects who attempted to try a lower or higher does if the original take-home does did not produce optimum results in combination with anti-nausea agents.
4. Subjects [and partners] who filled out and returned take-home questionnaires.

The practice of the present invention is demonstrated in the following examples. These examples are meant to illustrate the invention rather than to limit its scope. Variations in the treating compositions which do not adversely affect the effectiveness of the apomorphine or the antiemetic agent will be evident to one skilled in the art, and are within the scope of this invention.

**EXAMPLE 1: Apomorphine/Nicotine Combination By Wet Granulation Technique - Composition A**

Composition A Tablets were prepared from the ingredients listed in Table XXIII, below. Each ingredient was weighed as indicated and passed through a #35 mesh screen (sieve opening of about 0.51 mm) to ensure granulation. A solution containing the apomorphine HCL, the citric acid, half the acesulfame-K, half the peppermint flavor and half the chocolate flavor was prepared by dissolving the ingredients into a mixture of equal volumes of purified water and ethanol, USP. The solution was mixed until clear, and then absorbed into the listed amount of microcrystalline cellulose (Avicel 302). The resulting wet mass, which will be labelled "Part A," was mixed in a porcelain dish at room temperature (20°C) for 30 minutes, and then partially dried to obtain a solid mass. The mass was next granulated by screening through a #50 mesh (ASTM) (sieve opening of about 0.297 mm) stainless steel screen. The wet granules were dried at about 60°C to 70°C for about 1 to 1.5 hours. The resulting dried granules were then passed through a #35 mesh screen (sieve opening of about 0.51 mm).

Table XXIII: Apomorphine/Nicotine Combination Tablet Composition

	Ingredient	mg/tablet
	Apomorphine HCL	4.0
	Nicotine Base	1.0
5	Acesulfame-K	4.0
	Microcrystalline Cellulose	37.5
	Peppermint flavor	2.5
	Chocolate natural flavor	2.0
	Citric acid	3.0
10	Hydroxypropylmethylcellulose	13.0
	Mannitol	80.0
	Magnesium stearate	3.0
	TOTAL	150.0

15 Separately, nicotine was added to and blended with all the remaining ingredients except for the magnesium stearate. Specifically, the nicotine was added to the second half of the acesulfame-K, half the peppermint flavor, half the chocolate flavor, the

20 hydroxypropylmethylcellulose (methocel E4M, premium), and the mannitol. The resulting blend will be labelled "Part B." Parts A and B were then combined and mixed for about 5 minutes in a V-shaped blender. Next,

25 magnesium stearate was added to the blender and blending continued for about 2 minutes.

The final mix was removed from the blender and fed into a Stoke's single punch tablet press fitted with biconvex 5/16" diameter tooling for tablet preparation. Tablets were prepared at various compressional forces,

30 yielding tablets of different hardness. In general, the harder the tablet the slower the release of the active ingredients therefrom.

For additional discussion on methods for preparing sublingual apomorphine tablets see U.S. Patent

35 No. 5,624,677 to El-Rashidy et al., which is

incorporated here by reference to the extent that it is not inconsistent.

The dissolution of apomorphine and nicotine for Composition A Tablets was measured using a USP Type II apparatus (USP XXIII) stirred at 40 rpm. The dissolution medium was 500 ml of 10 millimolar ammonium phosphate buffer at a pH of  $3.0 \pm 0.5$  at about  $37^\circ\text{C}$ . The amount of apomorphine and nicotine released into the medium was detected by measuring absorbance at two different wavelengths, 259 nm and 272 nm, and resolving the following two equations:

$$A_{T259} = (\epsilon^{259}_{apo}) (C_{apo}) (l) + (\epsilon^{259}_{nic}) (C_{nic}) (l) \quad (5)$$

$$A_{T272} = (\epsilon^{272}_{apo}) (C_{apo}) (l) + (\epsilon^{272}_{nic}) (C_{nic}) (l) \quad (6)$$

In the above equations,  $A_{T259}$  is the total absorbance at 259 nanometers (nm);  $A_{T272}$  is the total absorbance at 272 nm;  $\epsilon^{259}_{apo}$  is the molar absorptivity of apomorphine at 259 nm;  $\epsilon^{259}_{nic}$  is the molar absorptivity of nicotine at 259 nm;  $\epsilon^{272}_{apo}$  is the molar absorptivity of apomorphine at 272 nm;  $\epsilon^{272}_{nic}$  is the molar absorptivity of nicotine at 272 nm;  $C_{apo}$  is the molar concentration of apomorphine;  $C_{nic}$  is the molar concentration of nicotine; and  $l$  is the cell path length.

By solving equations (5) and (6), the molar concentration of apomorphine ( $C_{apo}$ ) and nicotine ( $C_{nic}$ ) can be calculated from total absorbance data ( $A_{T259}$  and  $A_{T272}$ ) as follows.

$$C_{apo} = (\epsilon^{272}_{nic} A_{T259} - \epsilon^{259}_{nic} A_{T272}) / (\epsilon^{259}_{apo} \epsilon^{272}_{nic} - \epsilon^{272}_{apo} \epsilon^{259}_{nic}) \quad (7)$$

$$C_{nic} = (\epsilon^{272}_{apo} A_{T259} - \epsilon^{259}_{apo} A_{T272}) / (\epsilon^{272}_{apo} \epsilon^{259}_{nic} - \epsilon^{259}_{apo} \epsilon^{272}_{nic}) \quad (8)$$

5 Dissolution kinetic constants ( $K_{diss}$ ) for apomorphine and nicotine were calculated assuming zero-order release kinetics.

10 The tablets prepared were compared against a commercially available soluble apomorphine HCl tablet for dissolution characterization. The results are presented in Table XXVII (below) and in FIGURE 5. Specifically, the time to 50 percent drug release ( $T_{50}$ ) and 90 percent drug release ( $T_{90}$ ) for both apomorphine and nicotine are reported together with dissolution constants.

15 In addition, tablet hardness was measured using a Computest Tablet Hardness Tester. These results are also reported in Table XXVII.

20 Composition A Tablets released apomorphine relatively slower as compared to the release of the antiemetic agent, nicotine.

**EXAMPLE 2: Apomorphine/Nicotine Combination By Wet Granulation Technique - Composition B**

25 Composition B Tablets were prepared from the ingredients listed in Table XXIII (above). Each ingredient was weighed as indicated and passed through a #35 mesh screen (sieve opening of about 0.51 mm) to ensure granulation. Apomorphine HCL, the hydroxypropylmethyl cellulose, the citric acid, the acesulfame-K, the peppermint flavor, and the chocolate flavor were blended together with the indicated amount of microcrystalline cellulose using 25 percent ethanol in deionized water. The solution was mixed until clear, and then absorbed into half the listed amount of

35 microcrystalline cellulose (Avicel 302). The resulting

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wet mass (Part A) was mixed in a porcelain mortar at room temperature (20°C) for about 30 minutes, and then partially dried to obtain a single piece. The mass was granulated using a #35 mesh hand screen (sieve opening of about 0.51 mm). The wet granules were dried at about 60°C to 70°C for about 1 to 1.5 hours, and periodically mixed during the drying stage. The resulting dried granules were then passed through a #35 mesh hand screen (sieve opening of about 0.51 mm).

Separately, nicotine was added to and blended with the second half of the microcrystalline cellulose and the mannitol (Part B). Parts A and B were then combined and mixed for about 5 minutes in a V-shaped blender. Next, magnesium stearate was added to the blender, followed by continued blending for about 2 minutes.

The final mix was removed from the blender and compressed into tablets using a Stoke's single punch tablet press fitted with 5/16" diameter biconvex tooling. Tablets were prepared at various compressional forces, yielding tablets of different hardness.

Dissolution of apomorphine and nicotine for Composition B Tablets was measured and reported as described in Example 1. The results are presented in Table XXVII (below) and in FIGURE 6. Composition B Tablets released apomorphine relatively slower as compared to the release of nicotine.

**EXAMPLE 3: Apomorphine/Nicotine Layered Tablet Combination - Composition C**

The ingredients listed in TABLE XXIV (below) were used to prepare a layered tablet having a core portion containing apomorphine HCL and an outer layer containing the antiemetic agent nicotine. All

ingredients were first passed through a #35 mesh hand screen (sieve opening of about 0.51 mm).

**Table XXIV: Apomorphine/Nicotine Layered Tablet Composition**

5		<b>Ingredient</b>	<b>mg/tablet</b>
		<b>Tablet core:</b>	
		Apomorphine HCL	4.0
		Acesulfame-K	1.6
		Microcrystalline Cellulose	21.6
10		Peppermint flavor	1.0
		Chocolate flavor	0.8
		Citric acid	1.2
		Hydroxypropylmethylcellulose	4.0
		Mannitol	24.6
15		Magnesium stearate	1.2
		<b>Tablet outer layer:</b>	
		Nicotine base	1.0
		Acesulfame-K	0.4
		Microcrystalline Cellulose	36.6
20		Mannitol	47.0
		Magnesium stearate	1.0
		Hydroxypropylmethylcellulose	4.0
		<b>TOTAL</b>	<b>150.0</b>

25           The core portion was prepared by dry mixing apomorphine HCL, citric acid, peppermint flavor, chocolate flavor and acesulfame-K. The resulting mixture was blended in a V-shaped blender for about 5 minutes. Hydroxypropylmethylcellulose was then added and the blending continued for an additional 5 minutes. The microcrystalline cellulose was then added to the blender and mixing was continued for yet another 5 minutes. Next mannitol was added to the blender, followed by another 5 minute stage of blending.

30

35           Finally, magnesium stearate was added and blended in for about 2 minutes.



The resulting mixture was transferred to a Stoke's tablet press fitted with 7/32" diameter biconvex tooling to generate tablet cores with a hardness of about 3 kilopascals (Kp).

5           The outer antiemetic layer was prepared by mixing nicotine with the listed amount of microcrystalline cellulose in a porcelain mortar until the mixture became homogeneous. The homogeneous mixture was then transferred to a V-shaped blender, where the  
10       listed amounts of mannitol, hydroxypropylmethyl-cellulose, and acesulfame-K were blended in for about 5 minutes. Magnesium stearate was then added followed by an additional 2 minutes of blending.

15           A portion of the nicotine mixture was then transferred to the die of the Stoke's tablet press fitted with 5/16" biconvex tooling. Next an apomorphine tablet core discussed above was placed in the die and then covered with another portion of the nicotine mixture. The nicotine mixture and core portion were  
20       finally compressed together to form layered tablets.

25           Dissolution of apomorphine and nicotine for Composition C Tablets was measured and reported as described in Example 1. The results are presented in Table XXVII (below) and in FIGURE 7. As expected, Composition C Tablets released nicotine from their outer layer relatively sooner and faster than the apomorphine from the core portion.

30       **EXAMPLE 4:       Apomorphine/Prochlorperazine Combination  
                      By Wet Granulation Technique -  
                      Composition D**

35           Composition D Tablets were prepared from the ingredients listed in Table XXV (below). Each ingredient was weighed as indicated and passed through a #35 mesh screen (sieve opening of about 0.51 mm) to ensure granulation. A solution containing the

apomorphine HCL, acesulfame-K, peppermint flavor, chocolate flavor, and citric acid was prepared by dissolving these ingredients into a mixture of equal volumes of distilled water and ethanol. The solution was mixed until clear, and then absorbed into the listed amount of microcrystalline cellulose (Avicel 302) by further mixing over a stainless steel pan at room temperature (20°C) for about 30 minutes. The mixture was partially dried before granulating with a #60 mesh hand screen (sieve opening of about 0.25 mm).

**Table XXV: Apomorphine/Prochlorperazine Combination Tablet Composition**

Ingredient	mg/tablet
Apomorphine HCL	4.0
Prochlorperazine HCL	5.0
Acesulfame-K	4.0
Microcrystalline Cellulose	37.5
Peppermint flavor	2.5
Chocolate flavor	2.0
Citric acid	3.0
Hydroxypropylmethylcellulose	10.0
Mannitol	68.0
Sodium alginate	10.0
Magnesium stearate	3.0
TOTAL	150.0

The resulting granules were dried at about 60°C to 70°C for about 2 hours. The dried granules were then mixed in a porcelain mortar and passed through a #35 mesh hand screen (sieve opening of about 0.51 mm).

All remaining ingredients listed in Table XXV, except the magnesium stearate, were blended with the dry granules for about 5 minutes using a V-shaped blender. After 5 minutes of blending, magnesium stearate was added and the blending repeated for an additional 5

minutes. The resulting blend was compressed into tablets using the Stoke's tablet press fitted with 5/16" biconvex tooling.

Composition D Tablets were evaluated as described for Example 1, except that absorbance was measured at 254 nm rather than 259 nm. The results are presented in Table XXVII (below) and in FIGURE 8. Composition D Tablets released apomorphine relatively slower as compared to the release of prochlorperazine.

**EXAMPLE 5: Apomorphine/Prochlorperazine Combination  
By Wet Granulation Technique -  
Composition E**

Composition E Tablets were prepared from the ingredients listed in Table XXV (above). Each ingredient was weighed as indicated and passed through a #35 mesh screen (sieve opening of about 0.51 mm) to ensure granulation. Apomorphine HCL, the hydroxypropylmethyl cellulose, the sodium alginate, the citric acid, the acesulfame-K, the peppermint flavor, and the chocolate flavor were blended using 25 percent ethanol in deionized water. The resulting wet mass (Part A) was mixed in a porcelain mortar at room temperature (20°C) for about 30 minutes, and then partially dried to obtain a single piece. The resulting mass was granulated using a #35 mesh hand screen (sieve opening of about 0.51 mm). The wet granules were dried at about 60°C to 70°C for about 1 to 1.5 hours, and periodically mixed during the drying stage. The resulting dried granules were then passed through a #35 mesh hand screen (sieve opening of about 0.51 mm).

Separately, prochlorperazine was added to and blended with the mannitol (Part B). Parts A and B were then combined and mixed for about 5 minutes in a V-shaped blender. Next, magnesium stearate was added to

the blender, followed by continued blending for about 2 minutes.

The final mix was removed from the blender and compressed into tablets using a Stoke's single punch tablet press fitted with 5/16" diameter biconvex tooling. Tablets were prepared at various compressional forces, yielding tablets of different hardness.

Dissolution of apomorphine and prochlorperazine for Composition E Tablets was measured and reported as described in Example 1. The results are presented in Table XXVII (below) and in FIGURE 9. Composition E Tablets released apomorphine relatively slower as compared to the release of prochlorperazine.

**EXAMPLE 6:      Apomorphine/Prochlorperazine Layered  
Tablet Combination - Composition F**

Composition F Tablets were prepared according to the instructions presented in Example 3, except that the ingredients of Table XXVI (below) were used. Prochlorperazine was substituted for nicotine and the sodium alginate was added in the same step as the hydroxypropylmethylcellulose.

**Table XXVI: Apomorphine/Prochlorperazine  
Layered Tablet Composition**

	Ingredient	mg/tablet
	<b>Tablet core:</b>	
5	Apomorphine HCL	4.0
	Acesulfame-K	1.6
	Microcrystalline Cellulose	20.0
	Peppermint flavor	1.0
	Chocolate flavor	0.8
10	Citric acid	1.2
	Hydroxypropylmethylcellulose	5.0
	Mannitol	20.2
	Sodium alginate	5.0
	Magnesium stearate	1.2
15	<b>Tablet outer layer:</b>	
	Prochlorperazine	5.0
	Acesulfame-K	0.4
	Microcrystalline Cellulose	35.6
	Mannitol	46.0
20	Magnesium stearate	1.0
	Hydroxypropylmethylcellulose	2.0
	<b>TOTAL</b>	<b>150.0</b>

Composition F Tablets were evaluated as described for Example 1. The results are presented in Table XXVII (below) and in FIGURE 10. Composition F Tablets released the antiemetic prochlorperazine relatively faster than the apomorphine, as expected.

**EXAMPLE 7: Apomorphine/Prochlorperazine Layered  
Tablet Combination - Composition G**

The ingredients listed in Table XXVI (above) were used to prepare a layered tablet having a core portion containing apomorphine HCL and an outer layer containing the antiemetic agent prochlorperazine. Each ingredient was weighed as indicated and passed through a

#35 mesh screen (sieve opening of about 0.51 mm) to ensure granulation.

5 The core portion was prepared by dissolving the apomorphine HCL, acesulfame-K, peppermint flavor, chocolate flavor, and citric acid into a mixture of equal volumes of distilled water and ethanol. The solution was mixed until clear, and then absorbed into the listed amount of microcrystalline cellulose (Avicel 302) by further mixing over a stainless steel pan at 10 room temperature (20°C) for about 30 minutes. The mixture was partially dried before granulating with a #60 mesh hand screen.

15 The resulting granules were dried at about 60°C to 70°C for about 2 hours. The dried granules were then mixed in a porcelain mortar and passed through a #35 mesh hand screen (sieve opening of about 0.51 mm).

20 All remaining core ingredients listed in Table XXVI, except the magnesium stearate, were blended with the dry granules for about 5 minutes using a V-shaped blender. After 5 minutes of blending, magnesium stearate was added and the blending repeated for an additional 2 minutes. The resulting blend was compressed into 60 mg tablet cores using the Stoke's tablet press fitted with 7/32" biconvex tooling.

25 The outer antiemetic layer was prepared by dissolving the prochlorperazine and acesulfame-K into a mixture of equal volumes of distilled water and ethanol. The solution was mixed until clear, and then absorbed into the listed amount of microcrystalline cellulose by 30 mixing over a stainless steel pan at room temperature (20°C) for about 30 minutes. The mixture was partially dried before granulating with a #60 mesh hand screen.

The resulting granules were dried at about 60°C to 70°C for about 2 hours, mixed in a porcelain

TABLE XXVI

mortar, and passed through a #35 mesh hand screen (sieve opening of about 0.51 mm). The mannitol and the hydroxypropylmethyl cellulose were blended with the dry outer-layer granules for about 5 minutes using a V-shaped blender. After 5 minutes of blending, magnesium stearate was added and the blending repeated for an additional 2 minutes.

Layered tablets were then prepared by compressing the outer-coating granules around tablet cores as described in Example 3.

Dissolution of apomorphine and prochlorperazine for Composition G Tablets was measured and reported as described in Example 1. The results are presented in Table XXVII (below) and in FIGURE 11. As expected, Composition G Tablets released prochlorperazine from their outer layer relatively sooner and faster than the apomorphine, which escapes from the core portion.

#### **EXAMPLE 8: Comparison Of Drug Release Profiles**

The dissolution profile of a commercially available soluble apomorphine HCl tablet (Apomorphine HCl, 6 mg of Apomorphine HCL in a 60 mg tablet) was analyzed as described for Example 1. The results of this test are shown graphically in FIGURE 12, and listed in Table XXVII (below).

Also reported in Table XXVII are the time to 50 percent drug release ( $T_{50}$ ), the time to 90 percent drug release ( $T_{90}$ ), and the calculated dissolution constants of both the apomorphine and antiemetic agent (nicotine or prochlorperazine) for each example composition.

Table XXVII: Comparison Of Release Profiles And Tablet Hardness

Apomorphine HCL			Antiemetic Agent			r <sup>2</sup>	Hardness, k <sub>p</sub>
Comp.	T <sub>50</sub>	T <sub>90</sub>	K <sub>diss</sub>	T <sub>50</sub>	T <sub>90</sub>		
A	60	60	1.44	10	25	0.908	4.5
B	12	40	3.92	5	12	0.912	4.7
C	13	45	4.29	8	20	0.899	2.5/4.9
D	>90	>90	0.13	90	>90	0.945	6.5
E	22	40	2.76	16	27	0.944	4.2
F	15	60	3.42	7	30	0.956	2.5/4.8
G	80	>120	0.67	<10	40	0.932	3.5/5.5
Soluable Apo Tablet	13	30	3.82	---	---	0.909	4.2

These data demonstrate the ability of the present invention to release antiemetic agent relatively sooner and faster than the apomorphine. FIGURES 5 through 13 are graphs generated from the data presented in TABLE XXVII. Significantly, and well represented in graphical form, compositions according to the present invention also release apomorphine at an advantageously slower rate than that of the commercial sublingual table.

Figure 13 is a composite graph of the dissolution profiles for a commercially available apomorphine soluble tablet and a Composition G Tablet (Example 7). This graph well demonstrates the advantage of a layered, staggered-release tablet according to the present invention.

The foregoing discussion, examples, and the reported studies are intended as illustrative of the present invention and are not to be taken as limiting. Still other variants within the spirit and scope of this invention are possible and will readily present themselves to those skilled in the art.